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OM nucleic - nucleic search, using sw model

Run on: November 9, 2005, 18:15:28 ; Search time 497 Seconds
(without alignments)
366.068 Million cell updates/sec

Title: US-09-937-057-9

Perfect score: 22

Sequence: 1 tgactgtgaacgttatagatga 22

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 9794790 seqs, 413409567 residues

Total number of hits satisfying chosen parameters: 11332426

Minimum DB seq length: 0

Maximum DB seq length: 100

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications NA:**

- 1: /cgn2_6/ptodata/2/pubpna/US07_PUBCOMB.seq.*
- 2: /cgn2_6/ptodata/2/pubpna/PCT_NEW_PUB.seq.*
- 3: /cgn2_6/ptodata/2/pubpna/US06_NEW_PUB.seq.*
- 4: /cgn2_6/ptodata/2/pubpna/US06_PUBCOMB.seq.*
- 5: /cgn2_6/ptodata/2/pubpna/US07_NEW_PUB.seq.*
- 6: /cgn2_6/ptodata/2/pubpna/PCTUS_PUBCOMB.seq.*
- 7: /cgn2_6/ptodata/2/pubpna/US08_NEW_PUB.seq.*
- 8: /cgn2_6/ptodata/2/pubpna/US08_PUBCOMB.seq.*
- 9: /cgn2_6/ptodata/2/pubpna/US09A_PUBCOMB.seq.*
- 10: /cgn2_6/ptodata/2/pubpna/US09B_PUBCOMB.seq.*
- 11: /cgn2_6/ptodata/2/pubpna/US09C_PUBCOMB.seq.*
- 12: /cgn2_6/ptodata/2/pubpna/US09D_PUBCOMB.seq.*
- 13: /cgn2_6/ptodata/2/pubpna/US09E_PUBCOMB.seq.*
- 14: /cgn2_6/ptodata/2/pubpna/US10A_PUBCOMB.seq.*
- 15: /cgn2_6/ptodata/2/pubpna/US10B_PUBCOMB.seq.*
- 16: /cgn2_6/ptodata/2/pubpna/US10C_PUBCOMB.seq.*
- 17: /cgn2_6/ptodata/2/pubpna/US10D_PUBCOMB.seq.*
- 18: /cgn2_6/ptodata/2/pubpna/US10E_PUBCOMB.seq.*
- 19: /cgn2_6/ptodata/2/pubpna/US10F_PUBCOMB.seq.*
- 20: /cgn2_6/ptodata/2/pubpna/US10G_PUBCOMB.seq.*
- 21: /cgn2_6/ptodata/2/pubpna/US10H_PUBCOMB.seq.*
- 22: /cgn2_6/ptodata/2/pubpna/US10I_PUBCOMB.seq.*
- 23: /cgn2_6/ptodata/2/pubpna/US10J_PUBCOMB.seq.*
- 24: /cgn2_6/ptodata/2/pubpna/US10K_PUBCOMB.seq.*
- 25: /cgn2_6/ptodata/2/pubpna/US10L_PUBCOMB.seq.*
- 26: /cgn2_6/ptodata/2/pubpna/US11_NEW_PUB.seq.*
- 27: /cgn2_6/ptodata/2/pubpna/US60_NEW_PUB.seq.*
- 28: /cgn2_6/ptodata/2/pubpna/US60_PUBCOMB.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	22	100.0	22	9	US-09-967-881-9
2	20.4	92.7	22	9	US-09-791-500-5
3	20.4	92.7	22	9	US-09-791-500-6
4	20.4	92.7	22	9	US-09-770-943-2
5	20.4	92.7	22	10	US-09-848-986-10

6	20.4	92.7	22	10	US-09-848-986-11	Sequence 11, Appl
7	20.4	92.7	22	16	US-10-233-121A-10	Sequence 10, Appl
8	20.4	92.7	22	16	US-10-219-143-5	Sequence 5, Appl
9	20.4	92.7	22	16	US-10-219-143-6	Sequence 6, Appl
10	20.4	92.7	22	17	US-10-412-151-5	Sequence 5, Appl
11	20.4	92.7	22	17	US-10-412-151-6	Sequence 6, Appl
12	19.4	88.2	22	16	US-10-233-121A-11	Sequence 11, Appl
13	18.8	85.5	22	9	US-09-802-686-1	Sequence 1, Appl
14	18.8	85.5	22	9	US-09-802-686-4	Sequence 4, Appl
15	18.8	85.5	22	9	US-09-802-686-9	Sequence 9, Appl
16	18.8	85.5	22	9	US-09-802-685-1	Sequence 1, Appl
17	18.8	85.5	22	9	US-09-802-685-4	Sequence 4, Appl
18	18.8	85.5	22	9	US-09-802-685-9	Sequence 9, Appl
19	18.8	85.5	22	9	US-09-802-685-12	Sequence 12, Appl
20	18.8	85.5	22	9	US-09-791-500-1	Sequence 1, Appl
21	18.8	85.5	22	9	US-09-791-500-3	Sequence 3, Appl
22	18.8	85.5	22	9	US-09-791-500-8	Sequence 8, Appl
23	18.8	85.5	22	9	US-09-802-376-1	Sequence 1, Appl
24	18.8	85.5	22	9	US-09-802-376-4	Sequence 4, Appl
25	18.8	85.5	22	9	US-09-802-376-9	Sequence 9, Appl
26	18.8	85.5	22	9	US-09-802-376-10	Sequence 10, Appl
27	18.8	85.5	22	9	US-09-774-403A-1	Sequence 1, Appl
28	18.8	85.5	22	9	US-09-774-403A-3	Sequence 3, Appl
29	18.8	85.5	22	9	US-09-770-943-1	Sequence 1, Appl
30	18.8	85.5	22	9	US-09-770-943-3	Sequence 3, Appl
31	18.8	85.5	22	9	US-09-770-943-10	Sequence 10, Appl
32	18.8	85.5	22	9	US-09-802-370-1	Sequence 1, Appl
33	18.8	85.5	22	9	US-09-802-370-4	Sequence 4, Appl
34	18.8	85.5	22	9	US-09-802-445-1	Sequence 1, Appl
35	18.8	85.5	22	9	US-09-802-445-4	Sequence 4, Appl
36	18.8	85.5	22	9	US-09-820-484-1	Sequence 1, Appl
37	18.8	85.5	22	9	US-09-820-484-3	Sequence 3, Appl
38	18.8	85.5	22	9	US-09-820-484-7	Sequence 7, Appl
39	18.8	85.5	22	9	US-09-828-505-1	Sequence 1, Appl
40	18.8	85.5	22	9	US-09-828-505-4	Sequence 4, Appl
41	18.8	85.5	22	9	US-09-967-881-1	Sequence 1, Appl
42	18.8	85.5	22	9	US-09-967-881-2	Sequence 2, Appl
43	18.8	85.5	22	9	US-09-967-881-3	Sequence 3, Appl
44	18.8	85.5	22	10	US-09-927-422A-1	Sequence 1, Appl
45	18.8	85.5	22	10	US-09-927-422A-4	Sequence 4, Appl

ALIGNMENTS

RESULT 1
US-09-967-881-9
; Sequence 9, Application US/09967881
; Publication No. US20020192184A1
; GENERAL INFORMATION:
; APPLICANT: Assistance Publique - Hopitaux de Paris
; APPLICANT: Institut National de la Sante et de la Recherche M
; APPLICANT: Carpentier, Antoine
; TITLE OF INVENTION: Use of Stabilised Oligonucleotides for Preparing A Medicament wit
; FILE REFERENCE: 267/246 US
; CURRENT FILING DATE: 2001-09-28
; NUMBER OF SEQ ID NOS: 48
; SOFTWARE: Patent in version 3.1
; SEQ ID NO 9
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Oligodeoxynucleotide
US-09-967-881-9

Query Match 100.0%; Score 22; DB 9; Length 22;
Best Local Similarity 100.0%; Pred. No. 1.7;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TGACTGTGAACGTTATAGATGA 22

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Db      1  TGAAGTGAACGTTATAGATGA 22
|||||
RESULT 2
US-09-791-500-5
; Sequence 5, Application US/09791500
; Patent No. US20020042387A1
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Rachmilewitz, Daniel
; TITLE OF INVENTION: Method for Treating Inflammatory Bowel
; TITLE OF INVENTION: Disease and Other Forms of Gastrointestinal Inflammation.
; FILE REFERENCE: 6510-202US1
; CURRENT FILING DATE: 2001-02-22
; NUMBER OF SEQ ID NOS: 39
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 5
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic polynucleotide sequence
US-09-791-500-5
Query Match          92.7%; Score 20.4; DB 9; Length 22;
Best Local Similarity 95.5%; Pred. No. 9.9;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      1  TGAAGTGAACGTTATAGATGA 22
Db      1  TGAAGTGAACGTTATAGATGA 22
|||||

RESULT 3
US-09-791-500-6
; Sequence 6, Application US/09791500
; Patent No. US20020042387A1
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Rachmilewitz, Daniel
; TITLE OF INVENTION: Method for Treating Inflammatory Bowel
; TITLE OF INVENTION: Disease and Other Forms of Gastrointestinal Inflammation.
; FILE REFERENCE: 6510-202US1
; CURRENT FILING DATE: 2001-02-22
; NUMBER OF SEQ ID NOS: 39
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 6
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic polynucleotide sequence
US-09-791-500-6
Query Match          92.7%; Score 20.4; DB 9; Length 22;
Best Local Similarity 95.5%; Pred. No. 9.9;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      1  TGAAGTGAACGTTATAGATGA 22
Db      1  TGAAGTGAACGTTATAGATGA 22
|||||

RESULT 4
US-09-770-943-2
; Sequence 2, Application US/09770943
; Publication No. US20020086839A1
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Roman, Mark
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; TITLE OF INVENTION: Inhibitors of DNA Immunostimulatory
; TITLE OF INVENTION: Sequence Activity
; FILE REFERENCE: 6510-173US1
; CURRENT APPLICATION NUMBER: US/09/770,943
; CURRENT FILING DATE: 2001-01-26
; PRIOR APPLICATION NUMBER: 09/092,314
; PRIOR FILING DATE: 1998-06-05
; PRIOR APPLICATION NUMBER: 60/048,794
; PRIOR FILING DATE: 1997-06-06
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-09-770-943-2
Query Match          92.7%; Score 20.4; DB 9; Length 22;
Best Local Similarity 95.5%; Pred. No. 9.9;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      1  TGAAGTGAACGTTATAGATGA 22
Db      1  TGAAGTGAACGTTATAGATGA 22
|||||

RESULT 5
US-09-848-986-10
; Sequence 10, Application US/09848986
; Publication No. US20030176373A1
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Lois, Augusto F.
; APPLICANT: Takabayashi, Kenji
; TITLE OF INVENTION: Agents that Modulate DNA-PK Activity and
; TITLE OF INVENTION: Methods of Use Thereof
; FILE REFERENCE: 06510168US1
; CURRENT APPLICATION NUMBER: US/09/848,986
; CURRENT FILING DATE: 2001-05-03
; PRIOR APPLICATION NUMBER: us 60/262321
; PRIOR FILING DATE: 2001-01-17
; PRIOR APPLICATION NUMBER: us 60/202,274
; PRIOR FILING DATE: 2000-05-05
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 10
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: ISS-ODN
US-09-848-986-10
Query Match          92.7%; Score 20.4; DB 10; Length 22;
Best Local Similarity 95.5%; Pred. No. 9.9;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      1  TGAAGTGAACGTTATAGATGA 22
Db      1  TGAAGTGAACGTTATAGATGA 22
|||||

RESULT 6
US-09-848-986-11
; Sequence 11, Application US/09848986
; Publication No. US20030176373A1
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Lois, Augusto F.
; APPLICANT: Takabayashi, Kenji
; TITLE OF INVENTION: Agents that Modulate DNA-PK Activity and
```

; TITLE OF INVENTION: Methods of Use Thereof
; FILE REFERENCE: 06510168US1
; CURRENT APPLICATION NUMBER: US/09/848,986
; CURRENT FILING DATE: 2001-05-03
; PRIOR APPLICATION NUMBER: us 60/262321
; PRIOR FILING DATE: 2001-01-17
; PRIOR APPLICATION NUMBER: us 60/202,274
; PRIOR FILING DATE: 2000-05-05
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 11
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: methylated ISS-ODN
; NAME/KEY: modified base
; LOCATION: (11)...(11)
; OTHER INFORMATION: m5c
US-09-848-986-11

Query Match 92.7%; Score 20.4; DB 10; Length 22;
Best Local Similarity 95.5%; Pred. No. 9.9;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGACTGTGAACGTTATAGATGA 22
|||||
Db 1 TGACTGTGAACGTTAGAGATGA 22

RESULT 7

US-10-233-121A-10
; Sequence 10, Application US/10233121A
; Publication No. US20030125284A1
; GENERAL INFORMATION:
; APPLICANT: RAZ, EYAL
; APPLICANT: LOIS, AUGUSTO
; APPLICANT: TAKABAYASHI, KENJI

; TITLE OF INVENTION: AGENTS THAT MODULATE DNA-PK ACTIVITY AND
; TITLE OF INVENTION: METHODS OF USE THEREOF
; FILE REFERENCE: UCAL-168DIV
; CURRENT APPLICATION NUMBER: US/10/233,121A
; CURRENT FILING DATE: 2003-03-11
; PRIOR APPLICATION NUMBER: US 09/848,986
; PRIOR FILING DATE: 2001-05-04
; PRIOR APPLICATION NUMBER: US 60/202,274
; PRIOR FILING DATE: 2000-05-05
; PRIOR APPLICATION NUMBER: US 60/262,321
; PRIOR FILING DATE: 2001-01-17
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 10
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: phosphodiester or phosphorothioate oligonucleotide

Query Match 92.7%; Score 20.4; DB 16; Length 22;
Best Local Similarity 95.5%; Pred. No. 9.9;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGACTGTGAACGTTATAGATGA 22
|||||
Db 1 TGACTGTGAACGTTAGAGATGA 22

RESULT 8

US-10-219-143-5
; Sequence 5, Application US/10219143
; Publication No. US20030130217A1
; GENERAL INFORMATION:
; APPLICANT: RAZ, EYAL
; APPLICANT: RACHMILEWITZ, DANIEL
; TITLE OF INVENTION: Method for Treating Inflammatory Bowel
; FILE REFERENCE: UCAL-202CON
; CURRENT APPLICATION NUMBER: US/10/412,151
; CURRENT FILING DATE: 2003-04-11

; APPLICANT: RAZ, EYAL
; APPLICANT: RACHMILEWITZ, DANIEL
; TITLE OF INVENTION: Method for Treating Inflammatory Bowel
; TITLE OF INVENTION: Disease and Other Forms of Gastrointestinal Inflammation.
; FILE REFERENCE: 6510-202US1
; CURRENT APPLICATION NUMBER: US/10/219,143
; CURRENT FILING DATE: 2002-08-13
; PRIOR APPLICATION NUMBER: US/09/791,500
; PRIOR FILING DATE: 2001-02-22
; NUMBER OF SEQ ID NOS: 39
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 5
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic polynucleotide sequence
US-10-219-143-5

Query Match 92.7%; Score 20.4; DB 16; Length 22;
Best Local Similarity 95.5%; Pred. No. 9.9;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGACTGTGAACGTTATAGATGA 22
|||||
Db 1 TGACTGTGAACGTTAGAGATGA 22

RESULT 9

US-10-219-143-6
; Sequence 6, Application US/10219143
; Publication No. US20030130217A1
; GENERAL INFORMATION:
; APPLICANT: RAZ, EYAL
; APPLICANT: RACHMILEWITZ, DANIEL

; TITLE OF INVENTION: Method for Treating Inflammatory Bowel
; TITLE OF INVENTION: Disease and Other Forms of Gastrointestinal Inflammation.
; FILE REFERENCE: 6510-202US1
; CURRENT APPLICATION NUMBER: US/10/219,143
; CURRENT FILING DATE: 2002-08-13
; PRIOR APPLICATION NUMBER: US/09/791,500
; PRIOR FILING DATE: 2001-02-22
; NUMBER OF SEQ ID NOS: 39
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 6
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic polynucleotide sequence

Query Match 92.7%; Score 20.4; DB 16; Length 22;
Best Local Similarity 95.5%; Pred. No. 9.9;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGACTGTGAACGTTATAGATGA 22
|||||
Db 1 TGACTGTGAACGTTAGAGATGA 22

RESULT 10

US-10-412-151-5
; Sequence 5, Application US/10412151
; Publication No. US20030176389A1
; GENERAL INFORMATION:
; APPLICANT: RAZ, EYAL
; APPLICANT: RACHMILEWITZ, DANIEL
; TITLE OF INVENTION: Method for Treating Inflammatory Bowel
; FILE REFERENCE: UCAL-202CON
; CURRENT APPLICATION NUMBER: US/10/412,151
; CURRENT FILING DATE: 2003-04-11

; PRIOR APPLICATION NUMBER: 09/791,500
; PRIOR FILING DATE: 2001-02-22
; PRIOR APPLICATION NUMBER: 60/184,256
; PRIOR FILING DATE: 2000-02-23
; NUMBER OF SEQ ID NOS: 39
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 5
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic polynucleotide sequence
US-10-412-151-5

Query Match 92.7%; Score 20.4; DB 17; Length 22;
Best Local Similarity 95.5%; Pred. No. 9.9;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 TGACTGTGAACGTTATAGATGA 22
|||||
Db 1 TGACTGTGAACGTTAGAGATGA 22

RESULT 11
US-10-412-151-6
; Sequence 6, Application US/10412151
; Publication No. US20030176389A1
; GENERAL INFORMATION:
; APPLICANT: Raz, Eval
; APPLICANT: Rachmilewitz, Daniel
; TITLE OF INVENTION: Method for Treating Inflammatory Bowel
; FILE OF INVENTION: Disease and Other Forms of Gastrointestinal Inflammation.
; FILE REFERENCE: UCAL-202CON
; CURRENT APPLICATION NUMBER: US/10/412,151
; CURRENT FILING DATE: 2003-04-11
; PRIOR APPLICATION NUMBER: 09/791,500
; PRIOR FILING DATE: 2001-02-22
; PRIOR APPLICATION NUMBER: 60/184,256
; PRIOR FILING DATE: 2000-02-23
; NUMBER OF SEQ ID NOS: 39
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 6
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic polynucleotide sequence
US-10-412-151-6

Query Match 92.7%; Score 20.4; DB 17; Length 22;
Best Local Similarity 95.5%; Pred. No. 9.9;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 TGACTGTGAACGTTATAGATGA 22
|||||
Db 1 TGACTGTGAACGTTAGAGATGA 22

RESULT 12
US-10-233-121A-11
; Sequence 11, Application US/10233121A
; Publication No. US20030125284A1
; GENERAL INFORMATION:
; APPLICANT: RAZ, EVAL
; APPLICANT: LOIS, AUGUSTO
; APPLICANT: TAKABAYASHI, KENJI
; TITLE OF INVENTION: AGENTS THAT MODULATE DNA-PK ACTIVITY AND
; FILE OF INVENTION: METHODS OF USE THEREOF
; FILE REFERENCE: UCAL-168DIV
; CURRENT APPLICATION NUMBER: US/10/233,121A
; CURRENT FILING DATE: 2003-03-11
; PRIOR APPLICATION NUMBER: US 09/848,986
; PRIOR FILING DATE: 2001-05-04

; PRIOR APPLICATION NUMBER: US 60/202,274
; PRIOR FILING DATE: 2000-05-05
; PRIOR APPLICATION NUMBER: US 60/262,321
; PRIOR FILING DATE: 2001-01-17
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 11
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 11
; OTHER INFORMATION: n = 5-methylcytidine-phosphodiester or
; OTHER INFORMATION: 5-methylcytidine-phosphorothioate
US-10-233-121A-11

Query Match 88.2%; Score 19.4; DB 16; Length 22;
Best Local Similarity 90.9%; Pred. No. 30;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 TGACTGTGAACGTTATAGATGA 22
|||||
Db 1 TGACTGTGAANGTTAGACATGA 22

RESULT 13
US-09-802-686-1
; Sequence 1, Application US/09802686
; Patent No. US20010046967A1
; GENERAL INFORMATION:
; APPLICANT: Dynavax Technologies Corporation
; APPLICANT: Van Nest, Gary
; TITLE OF INVENTION: METHODS OF PREVENTING AND TREATING
; FILE OF INVENTION: RESPIRATORY VIRAL INFECTION USING IMMUNOMODULATORY
; FILE REFERENCE: POLYNUCLEOTIDE SEQUENCES
; FILE REFERENCE: 377882000900
; CURRENT APPLICATION NUMBER: US/09/802,686
; CURRENT FILING DATE: 2001-03-09
; PRIOR APPLICATION NUMBER: 60/188,583
; PRIOR FILING DATE: 2000-03-10
; NUMBER OF SEQ ID NOS: 10
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Polynucleotide containing CG
US-09-802-686-1

Query Match 85.5%; Score 18.8; DB 9; Length 22;
Best Local Similarity 90.9%; Pred. No. 57;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 TGACTGTGAACGTTATAGATGA 22
|||||
Db 1 TGACTGTGAACGTTTCGAGATGA 22

RESULT 14
US-09-802-686-4
; Sequence 4, Application US/09802686
; Patent No. US20010046967A1
; GENERAL INFORMATION:
; APPLICANT: Dynavax Technologies Corporation
; APPLICANT: Van Nest, Gary
; TITLE OF INVENTION: METHODS OF PREVENTING AND TREATING
; FILE OF INVENTION: RESPIRATORY VIRAL INFECTION USING IMMUNOMODULATORY
; FILE REFERENCE: POLYNUCLEOTIDE SEQUENCES
; FILE REFERENCE: 377882000900
; CURRENT APPLICATION NUMBER: US/09/802,686
; CURRENT FILING DATE: 2001-03-09

Mon Nov 14 10:15:28 2005

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; PRIOR APPLICATION NUMBER: 60/188,583
; PRIOR FILING DATE: 2000-03-10
; NUMBER OF SEQ ID NOS: 10
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 4
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Polynucleotide containing CG
US-09-802-686-4

Query Match      85.5%; Score 18.8; DB 9; Length 22;
Best Local Similarity 90.9%; Pred. No. 57;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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QY 1 TGACTGTGAACGTTATAGATGA 22
    |||||
DB 1 TGACTGTGAACGTTCCAGATGA 22
    |||||

```

```

RESULT 15
US-09-802-686-9
; Sequence 9, Application US/09802686
; Patent No. US20010046967A1
; GENERAL INFORMATION:
; APPLICANT: Dynavax Technologies Corporation
; APPLICANT: Van Nest, Gary
; TITLE OF INVENTION: METHODS OF PREVENTING AND TREATING
; TITLE OF INVENTION: RESPIRATORY VIRAL INFECTION USING IMMUNOMODULATORY
; TITLE OF INVENTION: POLYNUCLEOTIDE SEQUENCES
; FILE REFERENCE: 377882000900
; CURRENT APPLICATION NUMBER: US/09/802,686
; CURRENT FILING DATE: 2001-03-09
; PRIOR APPLICATION NUMBER: 60/188,583
; PRIOR FILING DATE: 2000-03-10
; NUMBER OF SEQ ID NOS: 10
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 9
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Polynucleotide not containing CG
US-09-802-686-9

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```

Query Match      85.5%; Score 18.8; DB 9; Length 22;
Best Local Similarity 90.9%; Pred. No. 57;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

```

```

QY 1 TGACTGTGAACGTTATAGATGA 22
    |||||
DB 1 TGACTGTGAAGGTTAGATGA 22
    |||||

```

Search completed: November 9, 2005, 19:28:46
Job time : 504 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: November 9, 2005, 18:04:54 ; Search time 1762 Seconds
(without alignments)
475.263 Million cell updates/sec

Title: US-09-937-057-9

Perfect score: 22

Sequence: 1 tgactgtgaacgttatgatga 22

Scoring table: IDENTITY NUC

Gapop 10.0, Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 675282

Minimum DB seq length: 0

Maximum DB seq length: 100

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:*

1: gb_est1:*

2: gb_est2:*

3: gb_hc:*

4: gb_est3:*

5: gb_est4:*

6: gb_est5:*

7: gb_est6:*

8: gb_gsl1:*

9: gb_gsl2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	15.6	70.9	81	9	CG620747
2	15.6	70.9	86	9	CG548982
3	15.6	70.9	90	9	CG546017
4	15.6	70.9	91	9	CG554726
5	15.6	70.9	97	9	CG631426
6	15.6	70.9	98	9	CG610654
7	15.6	70.9	100	9	CG496629
8	14.6	66.4	58	1	AI662714
9	14.6	66.4	58	8	BH908898
10	14.4	65.5	70	9	CG546804
11	14.4	65.5	80	6	CD384499
12	14.2	64.5	72	9	CG631425
13	14.2	64.5	97	4	BI881470
14	14	63.6	48	8	AZ843479
15	14	63.6	63	8	AZ431742
16	14	63.6	63	9	CG631431
17	14	63.6	67	8	BH911627
18	14	63.6	84	7	CK103587
19	14	63.6	86	9	CG640878
20	14	63.6	88	1	AI953694
21	14	63.6	90	9	AG195135
22	14	63.6	97	6	CD866080
23	14	63.6	98	1	AA464890
24	14	63.6	100	7	CV064730

25	13.6	61.8	62	1	AV403764
26	13.6	61.8	64	4	BG514647
27	13.6	61.8	78	9	CG562292
28	13.6	61.8	80	2	BB329424
29	13.6	61.8	81	7	CO514377
30	13.6	61.8	82	1	AA573133
31	13.6	61.8	92	1	AA462609
32	13.6	61.8	92	9	CG578258
33	13.6	61.8	100	1	AI971012
34	13.6	61.8	100	2	AW355356
35	13.6	61.8	100	7	H16183
36	13.4	60.9	51	9	AG199950
37	13.4	60.9	56	8	BZ664941
38	13.4	60.9	82	9	AG198560
39	13.4	60.9	88	8	AZ783178
40	13.2	60.0	40	1	AA779179
41	13.2	60.0	50	1	AI241767
42	13.2	60.0	68	9	CC797276
43	13.2	60.0	75	9	EX132834
44	13.2	60.0	79	7	CR560685
45	13.2	60.0	83	8	BZ765063

ALIGNMENTS

RESULT 1
CG620747
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
FEATURES
ORIGIN
Query Match
Best Local Similarity
Matches
QY

81 bp mRNA linear GSS 02-OCT-2003
OST318195 Mus musculus 129Sv/Ev Mus musculus cDNA clone OST18195,
mRNA sequence.
CG620747
CG620747.1 GI:37444596
GSS.
Mus musculus (house mouse)
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 81)
Zambrowicz, B.P., Abuin, A., Ramirez-Solis, R., Richter, L.J.,
Piggott, J., BeltrandelRio, H., Buxton, E.C., Edwards, J., Finch, R.A.,
Fridde, C.J., Gupta, A., Hansen, G., Hu, Y., Huang, W., Jaing, C.,
Key, B.W. Jr., Kipp, P., Kohlhauff, B., Ma, Z.-Q., Markesich, D.,
Payne, R., Potter, D.G., Qian, N., Shaw, J., Schrick, J., Shi, Z.-Z.,
Sparks, M.J., Van Sligtenhorst, I., Vogel, P., Walke, W., Xu, N.,
Zhu, Q., Person, C. and Sands, A.T.
Wnk1 kinase deficiency lowers blood pressure in mice: a gene-trap
screen to identify potential targets for therapeutic intervention
Proc. Natl. Acad. Sci. U.S.A. 100 (24), 14109-14114 (2003)
Contact: Zambrowicz BP
OmniBank
Lexicon Genetics Incorporated
4000 Research Forest Drive, The Woodlands, TX 77381, USA
Email: material@lexgen.com
Gene trap sequence tag generated by 3' RACE from mouse ES cells as
described in Zambrowicz et al (Nature. 1998 Apr 9;392(6676):608-11)
Class: Gene Trap.
Location/Qualifiers
1..81
/organism="Mus musculus"
/mol_type="mRNA"
/strain="129Sv/Ev"
/db_xref="taxon:10090"
/clone="OST318195"
/cell_type="embryonic stem cell"
/clone_lib="Mus musculus 129Sv/Ev"

70.9%; Score 15.6; DB 9; Length 81;
81.8%; Pred No. 5.7e+03;
Conservative 0; Mismatches 4; Indels 0; Gaps 0;
1 TGACTGTGACGTTATAGATGA 22


```

ORIGIN
Query Match          70.9%; Score 15.6; DB 9; Length 91;
Best Local Similarity 81.8%; Pred. No. 5.8e+03;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Oy 1 TGACTGTGAACGTTATAGATGA 22
||||| ||||| ||||| |||||
Db 40 TGACCGAGAACGTGATAGAGGA 61

RESULT 5
CG631426          97 bp mRNA linear GSS 02-OCT-2003
LOCUS OST347804 Mus musculus 129Sv/Ev Mus musculus cDNA clone OST347804,
DEFINITION mRNA sequence.
ACCESSION CG631426
VERSION GSS.
KEYWORDS GI:37455275
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 97)
AUTHORS Zambrowicz,B.P., Abuin,A., Ramirez-Solis,R., Richter,L.J.,
Piggott,J., BeltrandelRio,H., Buxton,E.C., Edwards,J., Finch,R.A.,
Friddle,C.J., Gupta,A., Hansen,G., Hu,Y., Huang,W., Jaing,C.,
Key,B.W. Jr., Kipp,P., Kohlhauff,B., Ma,Z.-Q., Markesich,D.,
Payne,R., Potter,D.G., Qian,N., Shaw,J., Schrick,J., Shi,Z.-Z.,
Sparks,M.J., Van Sligtenhorst,I., Vogel,P., Walke,W., Xu,N.,
Zhu,Q., Person,C. and Sands,A.T.
Wnk1 kinase deficiency lowers blood pressure in mice: a gene-trap
screen to identify potential targets for therapeutic intervention
Proc. Natl. Acad. Sci. U.S.A. 100 (24), 14109-14114 (2003)
Contact: Zambrowicz BP
OmniBank
Lexicon Genetics Incorporated
4000 Research Forest Drive, The Woodlands, TX 77381, USA
Email: materials@lexgen.com
Gene trap sequence tag generated by 3' RACE from mouse ES cells as
described in Zambrowicz et al (Nature. 1998 Apr 9;392(6676):608-11)
Class: Gene Trap.

FEATURES
Location/Qualifiers
source 1..98
/organism="Mus musculus"
/mol_type="mRNA"
/strain="129Sv/Ev"
/db_xref="taxon:10090"
/clone="OST294314"
/cell_type="embryonic stem cell"
/clone_lib="Mus musculus 129Sv/Ev"

ORIGIN
Query Match          70.9%; Score 15.6; DB 9; Length 98;
Best Local Similarity 81.8%; Pred. No. 5.8e+03;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Oy 1 TGACTGTGAACGTTATAGATGA 22
||||| ||||| ||||| |||||
Db 47 TGACCGAGAACGTGATAGAGGA 68

RESULT 7
CG496629          100 bp mRNA linear GSS 01-OCT-2003
LOCUS OST36752 Mus musculus 129Sv/Ev Mus musculus cDNA clone OST36752,
DEFINITION mRNA sequence.
ACCESSION CG496629
VERSION GSS.
KEYWORDS GI:37265188
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 100)
AUTHORS Zambrowicz,B.P., Abuin,A., Ramirez-Solis,R., Richter,L.J.,
Piggott,J., BeltrandelRio,H., Buxton,E.C., Edwards,J., Finch,R.A.,
Friddle,C.J., Gupta,A., Hansen,G., Hu,Y., Huang,W., Jaing,C.,
Key,B.W. Jr., Kipp,P., Kohlhauff,B., Ma,Z.-Q., Markesich,D.,
Payne,R., Potter,D.G., Qian,N., Shaw,J., Schrick,J., Shi,Z.-Z.,
Sparks,M.J., Van Sligtenhorst,I., Vogel,P., Walke,W., Xu,N.,
Zhu,Q., Person,C. and Sands,A.T.
Wnk1 kinase deficiency lowers blood pressure in mice: a gene-trap
screen to identify potential targets for therapeutic intervention
Proc. Natl. Acad. Sci. U.S.A. 100 (24), 14109-14114 (2003)
Contact: Zambrowicz BP
OmniBank
Lexicon Genetics Incorporated
4000 Research Forest Drive, The Woodlands, TX 77381, USA
Email: materials@lexgen.com
Gene trap sequence tag generated by 3' RACE from mouse ES cells as
described in Zambrowicz et al (Nature. 1998 Apr 9;392(6676):608-11)
Class: Gene Trap.

FEATURES
Location/Qualifiers
source 1..100
/organism="Mus musculus"
/mol_type="mRNA"
/strain="129Sv/Ev"
/db_xref="taxon:10090"
/clone="OST347804"
/cell_type="embryonic stem cell"
/clone_lib="Mus musculus 129Sv/Ev"

ORIGIN
Query Match          70.9%; Score 15.6; DB 9; Length 97;
Best Local Similarity 81.8%; Pred. No. 5.8e+03;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Oy 1 TGACTGTGAACGTTATAGATGA 22
||||| ||||| ||||| |||||
Db 46 TGACCGAGAACGTGATAGAGGA 67

RESULT 6
CG610654          98 bp mRNA linear GSS 02-OCT-2003
LOCUS OST294314 Mus musculus 129Sv/Ev Mus musculus cDNA clone OST294314,
DEFINITION mRNA sequence.
ACCESSION CG610654
VERSION GSS.
KEYWORDS GI:37434503
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

```

```

/organism="Mus musculus"
/mol_type="mRNA"
/strain="129Sv/Ev"
/db_xref="taxon:10090"
/clone="OST36752"
/cell_type="embryonic stem cell"
/clone_lib="Mus musculus 129Sv/Ev"

ORIGIN
Query Match      70.9%; Score 15.6; DB 9; Length 100;
Best Local Similarity 81.8%; Pred. No. 5.9e+03;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTATAGATGA 22
DB 49 TGACCGAAGACGTGATAGAGGA 70

RESULT 8
AI662714/c
LOCUS      58 bp      mRNA      linear      EST 10-MAY-1999
DEFINITION IMAGE:746496 3' similar to TR:088760 O88760 AP-9 PROTEIN. ;, mRNA
sequence.
ACCESSION AI662714
VERSION    AI662714.1 GI:4766297
KEYWORDS  EST.
SOURCE    Mus musculus (house mouse)
ORGANISM  Mus musculus

REFERENCE
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 58)
AUTHORS  Marra,M., Hillier,L., Kucaba,T., Martin,J., Beck,C., Wylie,T.,
Underwood,K., Steptoe,M., Theising,B., Allen,M., Bowers,Y.,
Person,B., Swaller,T., Gibbons,M., Pape,D., Harvey,N., Schurk,R.,
Ritter,E., Kohn,S., Shin,T., Jackson,Y., Cardenas,M., McCann,R.,
Waterston,K. and Wilson,R.
The WashU-NCI Mouse EST Project 1999
Unpublished (1999)
Other ESTs: va88c01.y1
Contact: Marra M/WashU-NCI Mouse EST Project 1999
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@watson.wustl.edu
This clone is available royalty-free through LNL; contact the
IMAGE Consortium (info@image.lnl.gov) for further information.
MGI:455480
This clone was previously sequenced on the 5' end only, this new
data is from the 3' end
Possible reversed clone: similarity on wrong strand
High quality sequence stop: 1.
FEATURES
Location/Qualifiers
1..58
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="IMAGE:746496"
/sex="unknown"
/tissue_type="fetus"
/dev_stage="12.5dpc total fetus"
/lab_host="DH10B"
/clone_lib="Soares mouse 3NME12 5"
/notes="Organ: whole fetus; Vector: pT7T3D-Pac (Pharmacia)
with a modified polylinker; Site_1: Not I; Site_2: Eco RI;
1st strand cDNA was primed with a Not I - oligo(dT) primer
15', TGTTACCAATCTGAAGTGGAGCGCCGCTATTTTITTTTTT
3', on total mouse RNA [provided by Minoru Ko, Wayne
State Univ.]; double-stranded cDNA was ligated to Eco RI
adaptors (Pharmacia), digested with Not I and cloned into
the Not I and Eco RI sites of the modified pT7T3 vector.

Library went through one round of normalization, and was
constructed by Bento Soares and M. Fatima Bonaldo. "

ORIGIN
Query Match      66.4%; Score 14.6; DB 1; Length 58;
Best Local Similarity 81.0%; Pred. No. 1.7e+04;
Matches 17; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 GACTGTGAACGTTATAGATGA 22
DB 29 GATTGTGACCTTATAGAGGA 9

RESULT 9
BH908898
LOCUS      58 bp      DNA      linear      GSS 04-SEP-2002
DEFINITION SALK_051118.17.90.x Arabidopsis thaliana TDNA insertion lines
Arabidopsis thaliana genomic clone SALK_051118.17.90.x, genomic
survey sequence.
ACCESSION BH908898
VERSION    BH908898.1 GI:22721831
KEYWORDS  GSS.
SOURCE    Arabidopsis thaliana (thale cress)
ORGANISM  Arabidopsis thaliana

REFERENCE
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsiis.
1 (bases 1 to 58)
AUTHORS  Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R.,
Gadrinab,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L.,
Shinn,P., Zimmerman,J. and Ecker,J.R.
A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
Unpublished (2001)
Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGNAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: ecker@salk.edu
This is single pass sequence recovered from the left border of
TDNA. This sequence lies within an annotated intron of Atlg1692.
Class: TDNA tagged.
FEATURES
Location/Qualifiers
1..58
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/db_xref="taxon:3702"
/clone_lib="Arabidopsis thaliana TDNA insertion lines"
/notes="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/tdna_protocols.html"

ORIGIN
Query Match      66.4%; Score 14.6; DB 8; Length 58;
Best Local Similarity 81.0%; Pred. No. 1.7e+04;
Matches 17; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTATAGATG 21
DB 9 TTACTGTTAAAGTAAATAGATG 29

RESULT 10
CG546804
LOCUS      70 bp      mRNA      linear      GSS 01-OCT-2003
DEFINITION OST146791 Mus musculus 129Sv/Ev Mus musculus cDNA clone OST146791,
CG546804/c

```

Diatom EST Database (<http://avesthagen.sznbowler.com>)

```

RESULT 13
BI881470          97 bp      mRNA      linear      EST 16-SEP-2002
LOCUS             Zbrafish Research Genetics C32 fin Danio rerio cDNA
DEFINITION        clone IMAGE:4468146 5' similar to TR:Q9W725 Q9W725 UNCOUPLING
SOURCE             PROTEIN 2. ;, mRNA sequence.
ORGANISM           BI881470.1 GI:16088741
VERSION            BI881470
KEYWORDS           Danio rerio (zebrafish)
SOURCE             Danio rerio
ORGANISM           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
                  Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
                  Cypriniformes; Cyprinidae; Danio.
REFERENCE          1 (bases 1 to 97)
AUTHORS            Clark,M., Johnson,S.L., Lehrach,H., Lee,R., Li,F., Marra,M.,
                  Eddy,S., Hillier,L., Kucaba,T., Martin,J., Beck,C., Wylie,T.,
                  Underwood,K., Steptoe,M., Theising,B., Allen,M., Bowers,Y.,
                  Person,B., Swaller,T., Gibbons,M., Pape,D., Harvey,N., Schurk,R.,
                  Ritter,E., Kohn,S., Shin,T., Jackson,Y., Cardenas,M., McCann,R.,
                  Waterston,R. and Wilson,R.
TITLE              WashU zebrafish EST Project 1998
JOURNAL            Unpublished (1998)
COMMENT            Contact: Stephen L. Johnson
                   Washington University School of Medicine
                   4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
                   Tel: 314 286 1800
                   Fax: 314 286 1810
                   Email: zbrafish@watson.wustl.edu
                   cDNA Library Preparation: Ning Wu. cDNA Library Arrayed by: Steve
                   Johnson. DNA Sequencing by: Washington University Genome Sequencing
                   Center NOTE: This clone is available royalty-free through LLNU;
                   contact the IMAGE Consortium (info.llnl.gov) for further
                   information.
FEATURES           Trace considered overall poor quality
Seq primer: T3 ET from Amersham
High quality sequence stop: 1.
Location/Qualifiers
    1..97
     /organism="Danio rerio"
     /mol_type="mRNA"
     /db_xref="taxon:7955"
     /clone="IMAGE:4468146"
     /tissue_type="Fin"
     /lab_host="GeneHogs (HS996, a phage-resistant isolate of
DH10B)"
     /clone_lib="Zebrafish Research Genetics C32 fin"
     /note="Vector: pTT73D-Pac with a modified polylinker;
Site_1: EcoRI; Site_2: NotI; let strand cDNA was prepared
from zebrafish(C32) fin, and was then primed with a Not I
- oligo(dT) primer. Double-stranded cDNA was ligated to
Eco RI adaptors (Pharmacia), digested with Not I and
cloned into the Not I and Eco RI sites of the modified
pTT73 vector. Library is non-normalized. Library was
constructed by Ning Wu. NOTE: This clone is available
royalty-free through LLNL; contact the IMAGE Consortium
(info.llnl.gov)for further information"
ORIGIN
Query Match      64.5%; Score 14.2; DB 4; Length 97;
Best Local Similarity 84.2%; Pred. No. 2,9e+04;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY   3 ACTGTGAACGTTATAGTG 21
      |||||
Db    38 ACCGTGAACGTTATTGGTG 56

RESULT 14
AZ843479          48 bp      DNA      linear      GSS 20-FEB-2001
LOCUS             2M0142C23F Mouse 10kb plasmid UUCG1M library Mus musculus genomic
DEFINITION        clone UUGC2M0142C23 F, genomic survey sequence.
SOURCE             AZ843479
VERSION            2M0142C23F
KEYWORDS           Mouse 10kb plasmid UUCG1M library Mus musculus genomic
SOURCE             clone UUGC2M0142C23 F, genomic survey sequence.
ORGANISM           AZ843479
VERSION            2M0142C23F
KEYWORDS           Mouse 10kb plasmid UUCG1M library Mus musculus genomic
SOURCE             clone UUGC2M0142C23 F, genomic survey sequence.
ORGANISM           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
                  Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE          1 (bases 1 to 48)
AUTHORS            Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
                  Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T.,
                  Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von
                  Niederhausern,A. and Wright,D., Weiss,R.
TITLE              Mouse whole genome scaffolding with paired end reads from 10kb
                  plasmid inserts
JOURNAL            Unpublished (2000)
COMMENT            Contact: Robert B. Weiss
                   University of Utah Genome Center
                   University of Utah
                   Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLCT, UT,
                   84112 USA
                   Tel: 801 585 5606
                   Fax: 801 585 7177
                   Email: ddunn@genetics.utah.edu
                   Insert Length: 10000 Std Error: 0.00
                   Plate: 0142 row: C column: 23
                   Seg primer: CGTTGTAAACGACGCCAGT
                   Class: plasmid ends
                   High quality sequence stop: 48.
FEATURES           Location/Qualifiers
    1..48
     /organism="Mus musculus"
     /mol_type="genomic DNA"
     /strain="C57BL/6J"
     /db_xref="taxon:10090"
     /clone="UUGC2M0142C23"
     /sex="Male"
     /lab_host="E. Coli strain XL10-Gold, TI-resistant, F-"
     /clone_lib="Mouse 10kb plasmid UUCG1M library"
     /note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pWD42 [gi|4732114|gb|AF129072.1], a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to
adaptor vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."
ORIGIN
Query Match      63.6%; Score 14; DB 8; Length 48;
Best Local Similarity 77.3%; Pred. No. 3.2e+04;
Matches 17; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY   1 TGACTGTGAACGTTATAGTGA 22
      |||||
Db    37 TGACTGTGTAATAAATAA 16

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LOCUS             1M0216018R Mouse 10kb plasmid UUCG1M library Mus musculus genomic
DEFINITION        clone UUGC1M0216018 R, genomic survey sequence.
SOURCE             AZ8431742
VERSION            1M0216018R
KEYWORDS           Mouse 10kb plasmid UUCG1M library Mus musculus genomic
SOURCE             clone UUGC1M0216018 R, genomic survey sequence.
ORGANISM           AZ8431742
VERSION            1M0216018R
KEYWORDS           Mouse 10kb plasmid UUCG1M library Mus musculus genomic
SOURCE             clone UUGC1M0216018 R, genomic survey sequence.
ORGANISM           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
                  Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE          1 (bases 1 to 48)
AUTHORS            Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
                  Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T.,
                  Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von
                  Niederhausern,A. and Wright,D., Weiss,R.
TITLE              Mouse whole genome scaffolding with paired end reads from 10kb
                  plasmid inserts
JOURNAL            Unpublished (2000)
COMMENT            Contact: Robert B. Weiss
                   University of Utah Genome Center
                   University of Utah
                   Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLCT, UT,
                   84112 USA
                   Tel: 801 585 5606
                   Fax: 801 585 7177
                   Email: ddunn@genetics.utah.edu
                   Insert Length: 10000 Std Error: 0.00
                   Plate: 0142 row: C column: 23
                   Seg primer: CGTTGTAAACGACGCCAGT
                   Class: plasmid ends
                   High quality sequence stop: 48.
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     /mol_type="genomic DNA"
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     /db_xref="taxon:10090"
     /clone="UUGC2M0142C23"
     /sex="Male"
     /lab_host="E. Coli strain XL10-Gold, TI-resistant, F-"
     /clone_lib="Mouse 10kb plasmid UUCG1M library"
     /note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pWD42 [gi|4732114|gb|AF129072.1], a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to
adaptor vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."
ORIGIN
Query Match      63.6%; Score 14; DB 8; Length 48;
Best Local Similarity 77.3%; Pred. No. 3.2e+04;
Matches 17; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY   1 TGACTGTGAACGTTATAGTGA 22
      |||||
Db    37 TGACTGTGTAATAAATAA 16

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ACCESSION AZ431742
VERSION AZ431742.1 GI:10555755
KEYWORDS GSS.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 63)
AUTHORS Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T.,
Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von
Niederhausern, A. and Wright, D., Weiss, R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 309, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0216 row: 0 column: 18
Seq primer: CACACAGGAACAGCTATGACC
Class: plasmid ends
High quality sequence stop: 63.
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/clone="UUC1M0216018"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, Tl-resistant, F-"
/clone_lib="Mouse 10kb plasmid UUC1M library"
/note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pWD42 (gi|4732114|gb|AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to
adaptor vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

ORIGIN
Query Match 63.6%; Score 14; DB 8; Length 63;
Best Local Similarity 77.3%; Pred. No. 3.4e+04;
Matches 17; Conservative 0; Mismatches 5; Indels 0; Gaps 0;
QY 1 TGACTGTGACGTATGATGA 22
||| ||||| ||| |||||
DB 36 TGAATGTGAATGTTTGAATGA 57

Search completed: November 9, 2005, 19:18:31
Job time : 1772 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: November 9, 2005, 18:06:40 ; Search time 94 Seconds
(without alignments)
382.958 Million cell updates/sec

Title: US-09-937-057-9

Perfect score: 22

Sequence: 1 tgactgtgaacgttatgatga 22

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Gapop 10.0 , Gapext 1.0

Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 1330268

Minimum DB seq length: 0

Maximum DB seq length: 100

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Issued Patents NA.*

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6: /cgn2_6/ptodata/1/ina/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	20.4	92.7	22	3	US-09-092-314-2
2	20.4	92.7	22	4	US-09-791-500-5
3	20.4	92.7	22	4	US-09-791-500-6
4	18.8	85.5	22	3	US-09-092-314-1
5	18.8	85.5	22	3	US-09-092-314-3
6	18.8	85.5	22	3	US-09-092-314-10
7	18.8	85.5	22	4	US-09-235-742-19
8	18.8	85.5	22	4	US-09-347-343-32
9	18.8	85.5	22	4	US-09-347-343-33
10	18.8	85.5	22	4	US-09-820-484-1
11	18.8	85.5	22	4	US-09-820-484-3
12	18.8	85.5	22	4	US-09-820-484-7
13	18.8	85.5	22	4	US-09-774-403A-1
14	18.8	85.5	22	4	US-09-774-403A-3
15	18.8	85.5	22	4	US-09-296-477-2
16	18.8	85.5	22	4	US-09-296-477-3
17	18.8	85.5	22	4	US-09-296-477-6
18	18.8	85.5	22	4	US-09-296-477-8
19	18.8	85.5	22	4	US-09-308-036A-1
20	18.8	85.5	22	4	US-09-791-500-1
21	18.8	85.5	22	4	US-09-791-500-3
22	18.8	85.5	22	4	US-09-791-500-8
23	18.8	85.5	22	4	US-09-565-906-2
24	18.4	83.6	22	4	US-09-296-477-16
25	17.8	80.9	22	4	US-09-296-477-12
26	17.8	80.9	22	4	US-09-296-477-15
27	17.2	78.2	22	3	US-09-092-314-4

28	17.2	78.2	22	4	US-09-235-742-20	Sequence 20, Appl
29	17.2	78.2	22	4	US-09-820-484-2	Sequence 2, Appl
30	17.2	78.2	22	4	US-09-820-484-6	Sequence 6, Appl
31	17.2	78.2	22	4	US-09-774-403A-2	Sequence 2, Appl
32	17.2	78.2	22	4	US-09-296-477-1	Sequence 1, Appl
33	17.2	78.2	22	4	US-09-296-477-5	Sequence 5, Appl
34	17.2	78.2	22	4	US-09-296-477-9	Sequence 9, Appl
35	17.2	78.2	22	4	US-09-296-477-13	Sequence 13, Appl
36	17.2	78.2	22	4	US-09-308-036A-2	Sequence 2, Appl
37	17.2	78.2	22	4	US-09-791-500-4	Sequence 4, Appl
38	17.2	78.2	22	4	US-09-791-500-9	Sequence 9, Appl
39	15.6	70.9	22	3	US-09-092-314-5	Sequence 5, Appl
40	15.6	70.9	22	3	US-09-092-314-7	Sequence 7, Appl
41	15.6	70.9	22	3	US-09-092-314-8	Sequence 8, Appl
42	15.6	70.9	22	4	US-09-791-500-2	Sequence 2, Appl
43	15.4	70.0	21	4	US-09-296-477-10	Sequence 10, Appl
44	15.4	70.0	23	4	US-09-296-477-11	Sequence 11, Appl
45	14.8	67.3	25	4	US-09-396-196G-55466	Sequence 55466, A

ALIGNMENTS

RESULT 1
US-09-092-314-2
; Sequence 2, Application US/09092314
; Patent No. 6225292
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Roman, Mark
; TITLE OF INVENTION: Inhibitors of DNA Immunostimulatory
; TITLE OF INVENTION: Sequence Activity
; Patent No. 6225292
; FILE REFERENCE: 6510-173US1
; CURRENT APPLICATION NUMBER: US/09/092,314
; CURRENT FILING DATE: 1998-06-05
; PRIOR APPLICATION NUMBER: 60/048,794
; PRIOR FILING DATE: 1997-06-06
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-09-092-314-2

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Qy 1 TGACTGTGAACGTTATAGTGA 22
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Db 1 TGACTGTGAACGTTATAGTGA 22

RESULT 2
US-09-791-500-5
; Sequence 5, Application US/09791500
; Patent No. 6613751
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Rachmilewitz, Daniel
; TITLE OF INVENTION: Method for Treating Inflammatory Bowel
; TITLE OF INVENTION: Disease and Other Forms of Gastrointestinal Inflammation.
; FILE REFERENCE: 6510-202US1
; CURRENT APPLICATION NUMBER: US/09/791,500
; CURRENT FILING DATE: 2001-02-22
; NUMBER OF SEQ ID NOS: 39
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 5
; LENGTH: 22

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; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic polynucleotide sequence
US-09-791-500-5

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Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db 1 TGACTGTGAACGTTAGAGATGA 22

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; Sequence 6, Application US/09791500
; Patent No. 6613751
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Rachmilewitz, Daniel
; TITLE OF INVENTION: Method for Treating Inflammatory Bowel
; TITLE OF INVENTION: Disease and Other Forms of Gastrointestinal Inflammation.
; FILE REFERENCE: 6510-202US1
; CURRENT APPLICATION NUMBER: US/09/791,500
; CURRENT FILING DATE: 2001-02-22
; NUMBER OF SEQ ID NOS: 39
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 6
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic polynucleotide sequence
US-09-791-500-6

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Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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US-09-092-314-1
; Sequence 1, Application US/09092314
; Patent No. 6225292
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Roman, Mark
; TITLE OF INVENTION: Inhibitors of DNA Immunostimulatory
; TITLE OF INVENTION: Sequence Activity
; Patent No. 6225292
; FILE REFERENCE: 6510-173US1
; CURRENT APPLICATION NUMBER: US/09/092,314
; CURRENT FILING DATE: 1998-06-05
; PRIOR APPLICATION NUMBER: 60/048,794
; PRIOR FILING DATE: 1997-06-06
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; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-09-092-314-1

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Best Local Similarity 90.9%; Pred. No. 6.3;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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; Sequence 3, Application US/09092314
; Patent No. 6225292
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Roman, Mark
; TITLE OF INVENTION: Inhibitors of DNA Immunostimulatory
; TITLE OF INVENTION: Sequence Activity
; Patent No. 6225292
; FILE REFERENCE: 6510-173US1
; CURRENT APPLICATION NUMBER: US/09/092,314
; CURRENT FILING DATE: 1998-06-05
; PRIOR APPLICATION NUMBER: 60/048,794
; PRIOR FILING DATE: 1997-06-06
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; OTHER INFORMATION: Oligonucleotide
US-09-092-314-3

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Db 1 TGACTGTGAACGTTAGAGATGA 22

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; Patent No. 6225292
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Roman, Mark
; TITLE OF INVENTION: Inhibitors of DNA Immunostimulatory
; TITLE OF INVENTION: Sequence Activity
; Patent No. 6225292
; FILE REFERENCE: 6510-173US1
; CURRENT APPLICATION NUMBER: US/09/092,314
; CURRENT FILING DATE: 1998-06-05
; PRIOR APPLICATION NUMBER: 60/048,794
; PRIOR FILING DATE: 1997-06-06
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 10
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-09-092-314-10

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; Patent No. 6498148
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; TITLE OF INVENTION: Immunization-Free Methods for Treating
; TITLE OF INVENTION: Antigen-Stimulated Inflammation in a Mammalian Host and
; TITLE OF INVENTION: Shifting the Host's Antigen Immune Responsiveness to a THI
; TITLE OF INVENTION: Phenotype
; FILE REFERENCE: 6510-170CON4
; CURRENT APPLICATION NUMBER: US/09/235.742
; CURRENT FILING DATE: 1999-01-21
; EARLIER APPLICATION NUMBER: 08/927,120
; EARLIER FILING DATE: 1997-09-05
; EARLIER APPLICATION NUMBER: 08/593,554
; EARLIER FILING DATE: 1996-01-30
; EARLIER APPLICATION NUMBER: 08/725,968
; EARLIER FILING DATE: 1996-10-04
; EARLIER APPLICATION NUMBER: 60/028,118
; EARLIER FILING DATE: 1996-10-11
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; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Recombinant or Synthetic Sequence
; US-09-235-742-19

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Best Local Similarity 90.9%; Pred. No. 6.3;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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      |||||
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RESULT 8
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; Patent No. 6514948
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal R.
; TITLE OF INVENTION: METHOD FOR ENHANCING AN IMMUNE RESPONSE
; FILE REFERENCE: 30448.64US01
; CURRENT APPLICATION NUMBER: US/09/347,343A
; CURRENT FILING DATE: 1999-07-02
; NUMBER OF SEQ ID NOS: 40
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 32
; LENGTH: 22
; TYPE: DNA
; ORGANISM: synthetic oligonucleotide
; US-09-347-343-32

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Best Local Similarity 90.9%; Pred. No. 6.3;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy      1 TGACTGTGAACGTTATAGATGA 22
      |||||
Db      1 TGACTGTGAACGTTTCGAGATGA 22

RESULT 9
US-09-347-343-33
; Sequence 33, Application US/09347343A
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; Patent No. 6514948
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal R.
; APPLICANT: KOBAYASHI, Hiroko
; TITLE OF INVENTION: METHOD FOR ENHANCING AN IMMUNE RESPONSE
; FILE REFERENCE: 30448.64US01
; CURRENT APPLICATION NUMBER: US/09/347,343A
; CURRENT FILING DATE: 1999-07-02
; NUMBER OF SEQ ID NOS: 40
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 33
; LENGTH: 22
; TYPE: DNA
; ORGANISM: synthetic oligonucleotide
; US-09-347-343-33

Query Match      85.5%; Score 18.8; DB 4; Length 22;
Best Local Similarity 90.9%; Pred. No. 6.3;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy      1 TGACTGTGAACGTTATAGATGA 22
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Db      1 TGACTGTGAACGTTTCGAGATGA 22

RESULT 10
US-09-820-484-1
; Sequence 1, Application US/09820484
; Patent No. 6534062
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Cho, Hearn Jay
; APPLICANT: Richman, Douglas
; APPLICANT: Horner, Anthony A.
; TITLE OF INVENTION: Method for Increasing a Cytotoxic T
; TITLE OF INVENTION: Lymphocyte Response in vivo.
; FILE REFERENCE: 06510-188US1
; CURRENT APPLICATION NUMBER: US/09/820,484
; CURRENT FILING DATE: 2001-03-28
; PRIOR APPLICATION NUMBER: US 60/192,537
; PRIOR FILING DATE: 2000-03-28
; PRIOR APPLICATION NUMBER: US 60/203,567
; PRIOR FILING DATE: 2000-05-11
; PRIOR APPLICATION NUMBER: US 60/215,895
; PRIOR FILING DATE: 2000-07-05
; NUMBER OF SEQ ID NOS: 8
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Disulfide-linked phosphorothioate ISS-ODN
; NAME/KEY: modified base
; LOCATION: (1)...(1)
; OTHER INFORMATION: disulfide thymine
; US-09-820-484-1

Query Match      85.5%; Score 18.8; DB 4; Length 22;
Best Local Similarity 90.9%; Pred. No. 6.3;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy      1 TGACTGTGAACGTTATAGATGA 22
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Db      1 TGACTGTGAACGTTTCGAGATGA 22

RESULT 11
US-09-820-484-3
; Sequence 3, Application US/09820484
; Patent No. 6534062
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
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; APPLICANT: Cho, Hearn Jay
; APPLICANT: Richman, Douglas
; APPLICANT: Horner, Anthony A.
; TITLE OF INVENTION: Method for Increasing a Cytotoxic T
; FILE REFERENCE: 06510-188US1
; CURRENT APPLICATION NUMBER: US/09/820,484
; CURRENT FILING DATE: 2001-03-28
; PRIOR APPLICATION NUMBER: US 60/192,537
; PRIOR FILING DATE: 2000-03-28
; PRIOR APPLICATION NUMBER: US 60/203,567
; PRIOR FILING DATE: 2000-05-11
; PRIOR APPLICATION NUMBER: US 60/215,895
; PRIOR FILING DATE: 2000-07-05
; NUMBER OF SEQ ID NOS: 8
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: phosphorothioate ISS-ODN
US-09-820-484-3

Query Match 85.5%; Score 18.8; DB 4; Length 22;
Best Local Similarity 90.9%; Pred. No. 6.3;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTATAGATGA 22
|||||
DB 1 TGACTGTGAACGTTTCGAGATGA 22

RESULT 12

US-09-820-484-7
; Sequence 7, Application US/09820484
; Patent No. 6534062
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Cho, Hearn Jay
; APPLICANT: Richman, Douglas
; APPLICANT: Horner, Anthony A.
; TITLE OF INVENTION: Method for Increasing a Cytotoxic T
; FILE REFERENCE: 06510-188US1
; CURRENT APPLICATION NUMBER: US/09/820,484
; CURRENT FILING DATE: 2001-03-28
; PRIOR APPLICATION NUMBER: US 60/192,537
; PRIOR FILING DATE: 2000-03-28
; PRIOR APPLICATION NUMBER: US 60/203,567
; PRIOR FILING DATE: 2000-05-11
; PRIOR APPLICATION NUMBER: US 60/215,895
; PRIOR FILING DATE: 2000-07-05
; NUMBER OF SEQ ID NOS: 8
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 7
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: MODN
US-09-820-484-7

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Best Local Similarity 90.9%; Pred. No. 6.3;
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DB 1 TGACTGTGAACCTTAGATGA 22

RESULT 13

US-09-774-403A-1
; Sequence 1, Application US/09774403A
; Patent No. 6552006
; GENERAL INFORMATION:
; APPLICANT: Eyal Raz
; APPLICANT: Richard Kornbluth
; APPLICANT: Antonio Catanzaro
; APPLICANT: Tomoko Hayashi
; APPLICANT: Dennis Carson
; TITLE OF INVENTION: Immunomodulatory Polynucleotides in
; FILE REFERENCE: UCAL166
; CURRENT APPLICATION NUMBER: US/09/774,403A
; CURRENT FILING DATE: 2002-04-15
; PRIOR APPLICATION NUMBER: 60/179,353
; PRIOR FILING DATE: 2000-01-31
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: FastSeq for Windows Version 4.0
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US-09-774-403A-1

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Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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RESULT 14

US-09-774-403A-3
; Sequence 3, Application US/09774403A
; Patent No. 6552006
; GENERAL INFORMATION:
; APPLICANT: Eyal Raz
; APPLICANT: Richard Kornbluth
; APPLICANT: Antonio Catanzaro
; APPLICANT: Tomoko Hayashi
; APPLICANT: Dennis Carson
; TITLE OF INVENTION: Immunomodulatory Polynucleotides in
; FILE REFERENCE: UCAL166
; CURRENT APPLICATION NUMBER: US/09/774,403A
; CURRENT FILING DATE: 2002-04-15
; PRIOR APPLICATION NUMBER: 60/179,353
; PRIOR FILING DATE: 2000-01-31
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US-09-774-403A-3

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US-09-296-477-2

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; Sequence 2, Application US/09296477A
; Patent No. 6589940
; GENERAL INFORMATION:
; APPLICANT: RAZ, E.
; APPLICANT: SCHWARTZ, D.
; APPLICANT: ROMAN, M.
; APPLICANT: DINA, D.
; TITLE OF INVENTION: IMMUNOSTIMULATORY OLIGONUCLEOTIDES,
; COMPOSITIONS THEREOF AND METHODS OF USE
; TITLE OF INVENTION: THEREOF
; FILE REFERENCE: 37782000420
; CURRENT APPLICATION NUMBER: US/09/296,477A
; CURRENT FILING DATE: 1999-04-22
; EARLIER APPLICATION NUMBER: 09/092,329
; EARLIER FILING DATE: 1998-06-05
; EARLIER APPLICATION NUMBER: 60/048,793
; EARLIER FILING DATE: 1997-06-06
; NUMBER OF SEQ ID NOS: 21
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US-09-296-477-2

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Job time : 96 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

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Run on: November 9, 2005, 17:34:52 ; Search time 1718 Seconds
(without alignments)
620.498 Million cell updates/sec

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Post-processing: Minimum Match 0%

Maximum Match 100%

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	22	100.0	22	6	AX036952
3	20.4	92.7	22	6	AR148608
4	20.4	92.7	22	6	BD136175
5	20.4	92.7	22	6	AR392166
6	20.4	92.7	22	6	AR392167
7	18.8	85.5	22	6	AR148607
8	18.8	85.5	22	6	AR148609
9	18.8	85.5	22	6	AR148616
10	18.8	85.5	22	6	BD136174
11	18.8	85.5	22	6	BD136176
12	18.8	85.5	22	6	BD136183
13	18.8	85.5	22	6	BD182369
14	18.8	85.5	22	6	BD185615
15	18.8	85.5	22	6	BD190435
16	18.8	85.5	22	6	BD190436
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19	18.8	85.5	22	6	BD233617

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22	18.8	85.5	22	6	BD233623	Immunost1
23	18.8	85.5	22	6	BD251283	Enhanceme
24	18.8	85.5	22	6	BD251284	Enhanceme
25	18.8	85.5	22	6	BD272056	Use of at
26	18.8	85.5	22	6	BD272057	Use of at
27	18.8	85.5	22	6	BD272058	Use of at
28	18.8	85.5	22	6	AR268334	Sequence
29	18.8	85.5	22	6	AR287741	Sequence
30	18.8	85.5	22	6	AR287743	Sequence
31	18.8	85.5	22	6	AR287745	Sequence
32	18.8	85.5	22	6	AR308057	Sequence
33	18.8	85.5	22	6	AR308059	Sequence
34	18.8	85.5	22	6	AR352573	Sequence
35	18.8	85.5	22	6	AR352574	Sequence
36	18.8	85.5	22	6	AR352577	Sequence
37	18.8	85.5	22	6	AR352579	Sequence
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40	18.8	85.5	22	6	AR392164	Sequence
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ACCESSION	BD272064.1				
VERSION	JP 2002539265-A/9				
KEYWORDS	synthetic construct				
SOURCE	synthetic construct				
ORGANISM	other sequences; artificial sequences.				
REFERENCE	1 (bases 1 to 22)				
AUTHORS	Carpentier, A.				
TITLE	Use of stabilized oligonucleotide for producing agents having antitumor activity				
JOURNAL	Patent: JP 2002539265-A 9 19-NOV-2002;				
COMMENT	ASSISTANCE PUBLIQUE HOPITAUX DE PARIS, INSTITUT NATIONAL DE LA SANTE ET DE LA RECHERCHE MEDICALE (INSERM)				
	OS Artificial Sequence				
	PN JP 2002539265-A/9				
	PD 19-NOV-2002				
	PF 17-MAR-2000 JP 2000606246				
	FR 19-MAR-1999 FR 99/03433				
	PI ANTOINE CARPENTIER				
	PC A61K47/48,A61K31/711,A61P35/00				
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METHOD for treating inflammatory bowel disease and other forms of
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JOURNAL Patent: US 6613751-A 6 02-SEP-2003;
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DEFINITION Sequence 1 from patent US 6225292.
ACCESSION ARI48607
VERSION ARI48607.1 GI:15112697
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 22)
Raz,E. and Roman,M.
TITLE Inhibitors of DNA immunostimulatory sequence activity
JOURNAL Patent: US 6225292-A 1 01-MAY-2001;
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Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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DEFINITION Sequence 3 from patent US 6225292.
ACCESSION ARI48609
VERSION ARI48609.1 GI:15112699
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 22)
Raz,E. and Roman,M.
TITLE Inhibitors of DNA immunostimulatory sequence activity
JOURNAL Patent: US 6225292-A 3 01-MAY-2001;
FEATURES Location/Qualifiers
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DEFINITION Sequence 10 from patent US 6225292.
ACCESSION ARI48616
VERSION ARI48616.1 GI:15112706
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 22)
Raz,E. and Roman,M.
TITLE Inhibitors of DNA immunostimulatory sequence activity
JOURNAL Patent: US 6225292-A 10 01-MAY-2001;
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|||||
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DEFINITION Inhibitors of DNA immunostimulatory sequence activity.
ACCESSION BD136174
VERSION BD136174.1 GI:23231119
KEYWORDS JP 2002505580-A/1.
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE 1 (bases 1 to 22)
Raz,E. and Roman,M.
TITLE Inhibitors of DNA immunostimulatory sequence activity
JOURNAL Patent: JP 2002505580-A 1 19-FEB-2002;
COMMENT DYNAVAX TECHNOLOGIES CORP, THE REGENTS OF THE UNIVERSITY OF CALIFORNIA
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PN JP 2002505580-A/1
PD 19-FEB-2002
PF 05-JUN-1998 JP 1999502803
PR 06-JUN-1997 US 60/048793
PI EYAL RAZ, MARK ROMAN
PC C12N15/00, C12N15/63, C12N15/79, C12N15/09, A61K48/00 CC
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ACCESSION  BD136176
VERSION     BD136176.1 GI:23231121
KEYWORDS   JP 2002505580-A/3
SOURCE     synthetic construct
ORGANISM   other sequences; artificial sequences.
REFERENCE  1 (bases 1 to 22)
AUTHORS    Raz,E. and Roman,M.
TITLE      Inhibitors of DNA immunostimulatory sequence activity
JOURNAL    Patent: JP 2002505580-A 3 19-FEB-2002;
DYNAXVAX TECHNOLOGIES CORP., THE REGENTS OF THE UNIVERSITY OF
CALIFORNIA
COMMENT    OS Artificial Sequence
PN JP 2002505580-A/3
PD 19-FEB-2002
PP 05-JUN-1998 JP 1999502803
PR 08-JUN-1997 US 60/048793
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DEFINITION Inhibitors of DNA immunostimulatory sequence activity.
ACCESSION  BD136183
VERSION     BD136183.1 GI:23231128
KEYWORDS   JP 2002505580-A/10
SOURCE     synthetic construct
ORGANISM   other sequences; artificial sequences.
REFERENCE  1 (bases 1 to 22)
AUTHORS    Raz,E. and Roman,M.
TITLE      Inhibitors of DNA immunostimulatory sequence activity
JOURNAL    Patent: JP 2002505580-A 10 19-FEB-2002;
DYNAXVAX TECHNOLOGIES CORP., THE REGENTS OF THE UNIVERSITY OF
CALIFORNIA
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PD 19-FEB-2002
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DEFINITION Anti-tumor antigens or their epitopes against HTLV-1 tumor.
ACCESSION  BD182369
VERSION     BD182369.1 GI:30793287
KEYWORDS   WO 02090981-A/1.
SOURCE     synthetic construct
ORGANISM   other sequences; artificial sequences.
REFERENCE  1 (bases 1 to 22)
AUTHORS    Hanabuchi,S., Ohashi,T. and Kannagi,M.
TITLE      Anti-tumor antigens or their epitopes against HTLV-1 tumor
JOURNAL    Patent: WO 02090981-A 1 14-NOV-2002;
JAPAN SCIENCE AND TECHNOLOGY CORP,SHINO HANABUCHI,TAKAKASHI OHASHI,
MARI KANNAGI
COMMENT    OS Artificial Sequence
PN WO 02090981-A/1
PD 14-NOV-2002
PP 02-MAY-2002 WO 2002JP004406
PR 08-MAY-2001 JP 01P 137526
PI SHINO HANABUCHI,TAKASHI OHASHI,MARI KANNAGI
PC G01N33/50,G01N33/15,A61K39/00
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DEFINITION Anti-tumor antigens or their epitopes against HTLV-I tumor.
ACCESSION  BD185615
VERSION     BD185615.1 GI:31877815
KEYWORDS   JP 2002372532-A/1.
SOURCE     synthetic construct
ORGANISM   other sequences; artificial sequences.
REFERENCE  1 (bases 1 to 22)
AUTHORS    Hanabuchi,S., Ohashi,T. and Kannagi,M.
TITLE      Anti-tumor antigens or their epitopes against HTLV-I tumor

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JAPAN SCIENCE AND TECHNOLOGY CORP
COMMENT OS Artificial Sequence
PN JP 2002372532-A/1
PD 26-DEC-2002
PF 08-MAY-2001 JP 2001137526
PI SHINO HANABUCHI, TAKASHI OHASHI, MARI KANNAGI
PC G01N33/50, A61K39/00, A61K39/21, A61P35/02, A61P37/04,
C07K1/06,
PC C12N5/06, C12Q1/02, G01N33/00, G01N33/15, G01N33/53, G01N33/53, PC
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DEFINITION BD190435
ACCESSION BD190435
VERSION BD190435.1 GI:33000174
KEYWORDS JP 2002537102-A/19.
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.
REFERENCE 1 (bases 1 to 22)
AUTHORS Barackman, J., Simph, M., Ugozoli, M., Kazazu, J., Donnelly, J.,
Ott, G.S. and Ohagan, D.
TITLE Microemulsions with Adsorbed Macromolecules and Microparticles
JOURNAL Patent: JP 2002537102-A 19 05-NOV-2002;
Chiron Corporation
COMMENT OS Artificial Sequence
PN JP 2002537102-A/19
PD 05-NOV-2002
PF 03-FEB-2000 JP 2000600618
PR 29-JUL-1999 US 60/146391, 28-OCT-1999 US 60/161997, PR
26-FEB-1999 US 60/121858
PI John Barackman, manmohan simph, mildred ugozoli, jina kazazu, john
PI donnelly,
PI Gary S Ott, Derek Ohagan
CC Oligonucleotide
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Copyright (c) 1993 - 2005 CompuGen Ltd.

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- 13: Geneseq2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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4	20.4	92.7	22	6	AAD24895 Methylate
5	20.4	92.7	22	6	AAD24894 Immunosti
6	20.4	92.7	22	12	ADO5351 Immune mo
7	19.4	88.2	22	12	ADO5287 Immune mo
8	18.8	85.5	22	2	AAV32079 Nucleotid
9	18.8	85.5	22	2	AAV80099 Immunomod
10	18.8	85.5	22	2	AAV80097 Immunomod
11	18.8	85.5	22	2	AAV80103 Immunomod
12	18.8	85.5	22	2	AAV80106 Oligo use
13	18.8	85.5	22	2	AAV80101 Immunomod
14	18.8	85.5	22	2	AAV80104 Oligo use
15	18.8	85.5	22	2	AAV80102 Immunomod
16	18.8	85.5	22	2	AAV55797 Immunosti
17	18.8	85.5	22	2	AAV55788 Immunosti
18	18.8	85.5	22	2	AAV55790 Immunosti
19	18.8	85.5	22	2	AAV36624 ISS-ODN D
20	18.8	85.5	22	3	AA14469 Mutant im

21	18.8	85.5	22	3	AA14467	AA14467 Immunosti
22	18.8	85.5	22	3	AA38072	AA38072 Immunosti
23	18.8	85.5	22	3	AA38070	AA38070 Immunosti
24	18.8	85.5	22	3	AA38068	AA38068 Immunosti
25	18.8	85.5	22	3	AA38071	AA38071 Immunosti
26	18.8	85.5	22	3	AA38065	AA38065 Immunosti
27	18.8	85.5	22	3	AA90459	AA90459 CpG adjuv
28	18.8	85.5	22	3	AA90458	AA90458 CpG adjuv
29	18.8	85.5	22	3	AA96254	AA96254 Sequence
30	18.8	85.5	22	3	AA96253	AA96253 Sequence
31	18.8	85.5	22	3	AA96252	AA96252 Sequence
32	18.8	85.5	22	3	AAZ55876	AAZ55876 Immunomod
33	18.8	85.5	22	3	AAC64052	AAC64052 Non-CpG c
34	18.8	85.5	22	3	AAC64051	AAC64051 Immunosti
35	18.8	85.5	22	4	AAH20404	AAH20404 CpG motif
36	18.8	85.5	22	4	AAH20403	AAH20403 CpG motif
37	18.8	85.5	22	4	AAH43345	AAH43345 Immunomod
38	18.8	85.5	22	4	AAH43340	AAH43340 Immunomod
39	18.8	85.5	22	4	AAH43338	AAH43338 Immunomod
40	18.8	85.5	22	4	AAH73439	AAH73439 Immunomod
41	18.8	85.5	22	4	AAH73441	AAH73441 Immunomod
42	18.8	85.5	22	4	AAH75992	AAH75992 Immunomod
43	18.8	85.5	22	4	AAI64301	AAI64301 Control o
44	18.8	85.5	22	4	AAH75995	AAH75995 Immunomod
45	18.8	85.5	22	4	AAH76000	AAH76000 Control o

ALIGNMENTS

RESULT 1

AAA96260
ID AAA96260 standard; DNA; 22 BP.

AC AAA96260;

DT 08-FEB-2001 (first entry)

DE Sequence of a stabilised oligonucleotide with antitumour activity.

KW Antitumour; immunostimulatory oligonucleotide; tumour; anaplasia;
KW Glioblastoma; medullablastoma; neuroblastoma; carcinoma; ss.

OS Synthetic.

PN WO200056342-A2.

PD 28-SEP-2000.

PF 17-MAR-2000; 2000WO-FR000676.

PR 19-MAR-1999; 99FR-00003433.

PA (ASSI-) ASSISTANCE PUBLIQUE HOPITAUX PARIS.
(INRM) INST NAT SANTE & RECH MEDICALE.

PI Carpentier A;

DR WPI; 2000-602192/57.

PT Use of stabilised oligonucleotides as antitumor agents, particularly
against nervous system tumors, have optimal activity and are not toxic.

XX Claim 3; Page 48; 57pp; French.

XX The present sequence represents a stabilised oligonucleotide which has
antitumour activity. The oligonucleotide comprises an octamer motif of
the type 5'-purine-purine-CG-pyrimidine-pyrimidine-X-X-3', where the pair
X-X is AT, AA, CT or TT. The oligonucleotides are immunostimulatory, and
are not toxic. They may be adapted for use in animals or humans. The
stabilised oligonucleotides are used for treating tumours, of any type
and any degree of anaplasia, particularly human tumours in the peripheral
or central nervous systems, specifically glioblastomas, medullablastomas,

```
CC neuroblastomas, melanomas or carcinomas
XX
SQ Sequence 22 BP; 7 A; 2 C; 6 G; 7 T; 0 U; 0 Other;

  Query Match      100.0%; Score 22; DB 3; Length 22;
  Best Local Similarity 100.0%; Pred. No. 0.65;
  Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTATAGATGA 22
   |||||
Db 1 TGACTGTGAACGTTATAGATGA 22
   |||||

RESULT 2
AAH43343
ID AAH43343 standard; DNA; 22 BP.
XX
AC AAH43343;
XX
DT 13-DEC-2001 (first entry)
XX
Immunomodulatory polynucleotide 1039.
DE
XX Immunomodulation; inflammation; gastrointestinal tract;
KW ulcerative colitis; Crohn's disease; inflammatory bowel disease;
KW diarrhoea; rectal bleeding; weight loss; colon; weight; lesion; ss.
XX
OS Synthetic.
XX
XX WO200162207-A2.
XX
XX 30-AUG-2001.
XX
XX 22-FEB-2001; 2001WO-US006034.
XX
XX 23-FEB-2000; 2000US-0184256P.
XX
XX (REGC ) UNIV CALIFORNIA.
XX
XX PI Raz E, Rachmilewitz D;
XX
XX WPI; 2001-565393/63.
XX
XX
XX Ameliorating gastrointestinal inflammation e.g. inflammatory bowel
PT disease involves administering an immunomodulatory nucleic acid.
XX
XX Claim 7; Page 28; 58pp; English.
XX
XX The sequences given in AAH43338-48 represent immunomodulatory
CC polynucleotides which may be used to ameliorate inflammation of the
CC gastrointestinal tract by administering a nucleic acid comprising one of
CC these sequences. These polynucleotides all comprise an immunomodulatory
CC nucleotide sequence of 5'-CpG-3' (I). The nucleotides may be used for
CC ameliorating or reducing gastrointestinal inflammation e.g. chronic or
CC acute gastrointestinal inflammation, ulcerative colitis, Crohn's disease
CC caused by inflammatory bowel disease, diarrhoea, rectal bleeding, weight
CC loss; to reduce colon weight and colon lesions; to reduce a colonic
CC inflammation. The immunomodulatory polynucleotides treat inflammatory
CC bowel disease satisfactorily and effectively and have little or no
CC toxicity even at a high dosage of 50000 micro-g. They also reduce the
CC risk of colonic cancer by treating ulcerative colitis
XX
XX Sequence 22 BP; 7 A; 2 C; 7 G; 6 T; 0 U; 0 Other;
SQ
  Query Match      92.7%; Score 20.4; DB 4; Length 22;
  Best Local Similarity 95.5%; Pred. No. 3.9;
  Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTATAGATGA 22
   |||||
Db 1 TGACTGTGAACGTTAGATGA 22
   |||||

RESULT 4
AAD24895
ID AAD24895 standard; DNA; 22 BP.
XX
AC AAD24895;
XX
DT 12-MAR-2002 (first entry)
XX
DE Methylated (5-methyl C) immunostimulatory oligodeoxynucleotide (ISS-ODN).
KW Cell death; DNA damage; DNA-dependent protein kinase; DNA-PK; necrosis;
KW immune response; apoptosis; Alzheimer's disease; Parkinson's disease;
KW rheumatoid arthritis; inflammation; osteoporosis; myocardial infarction;
```

KW liver disease; reperfusion injury; carcinoma; multiple sclerosis; stroke;
KW amyotrophic lateral sclerosis; Acquired Immune Deficiency Syndrome; AIDS;
KW head injury damage; aplastic anaemia; tumour; organ transplantation;
KW cerebral infarction; follicular lymphomas; systemic lupus erythematosus;
KW viral infection; glomerulonephritis; apoptosis; autoimmune disorder;
KW sepsis; immunostimulatory oligodeoxynucleotide; ISS-ODN; ss.

XX Unidentified.

XX Key Location/Qualifiers

FT modified_base 11

FT /*tag= a

FT /mod_base= m5c

XX WO200185910-A2.

XX 15-NOV-2001.

XX 04-MAY-2001; 2001WO-US014508.

XX 05-MAY-2000; 2000US-0202274P.

XX 17-JAN-2001; 2001US-0262321P.

XX (REGC) UNIV CALIFORNIA.

XX Raz E, Lois AF, Takabayashi K;

XX WPI; 2002-062244/08.

XX Modulating cell death or reducing DNA damage in eukaryotic cells, useful
PT for reducing cell death in individual or organ, comprises contacting cell
PT with agent modulating biological activity of DNA-dependent protein
PT kinase.

XX Example 3; Page 33; 57pp; English.

XX The invention relates to a method for modulating cell death or reducing
CC DNA damage in an eukaryotic cell by contacting the cell with an agent
CC that modulates the biological activity of DNA-dependent protein kinase
CC (DNA-PK). The invention also relates nucleic acids which modulate the
CC immune response binding to Ku antigen, resulting in activation of DNA-PK.
CC The method is useful for modulating cell death or reducing DNA damage in
CC an eukaryotic cell, for treating any disorder resulting from a genotoxic
CC insert to a cell e.g., necrosis, apoptosis. The method is also useful for
CC treating cell death-related indications such as Alzheimer's disease,
CC Parkinson's disease, rheumatoid arthritis, septic shock, sepsis, stroke,
CC central nervous system inflammation, osteoporosis, degenerative liver
CC disease, cerebellar degeneration, reperfusion injury, multiple sclerosis,
CC amyotrophic lateral sclerosis, myocardial infarction, head injury damage,
CC acquired immunodeficiency syndrome (AIDS), aplastic anaemia, cerebral
CC infarction, bypass heart surgery, organ transplantation. The method is
CC also useful for treating follicular lymphomas, carcinomas, autoimmune
CC disorders (systemic lupus erythematosus), hormone dependent tumours, The
CC immune mediated glomerulonephritis; apoptosis and viral infections. The
CC present sequence is methylated (5-methyl C) immunostimulatory
CC oligodeoxynucleotide (ISS-ODN) used in the exemplification of the
CC invention

XX Sequence 22 BP; 7 A; 2 C; 7 G; 6 T; 0 U; 0 Other;

Query Match 92.7%; Score 20.4; DB 6; Length 22;
Best Local Similarity 95.5%; Pred. No. 3.9;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTATAGATGA 22

DB 1 TGACTGTGAACGTTATAGATGA 22

RESULT 5

AAD24894

ID AAD24894 standard; DNA; 22 BP.

XX

AC AAD24894;

XX 12-MAR-2002 (first entry)

XX Immunostimulatory oligodeoxynucleotide (ISS-ODN) 2.

XX Cell death; DNA damage; DNA-dependent protein kinase; DNA-PK; necrosis;
KW immune response; apoptosis; Alzheimer's disease; Parkinson's disease;
KW rheumatoid arthritis; inflammation; osteoporosis; myocardial infarction;
KW liver disease; reperfusion injury; carcinoma; multiple sclerosis; stroke;
KW amyotrophic lateral sclerosis; Acquired Immune Deficiency Syndrome; AIDS;
KW head injury damage; aplastic anaemia; tumour; organ transplantation;
KW cerebral infarction; follicular lymphomas; systemic lupus erythematosus;
KW viral infection; glomerulonephritis; apoptosis; autoimmune disorder;
KW sepsis; immunostimulatory oligodeoxynucleotide; ISS-ODN; ss.

XX Unidentified.

XX WO200185910-A2.

XX 15-NOV-2001.

XX 04-MAY-2001; 2001WO-US014508.

XX 05-MAY-2000; 2000US-0202274P.

XX 17-JAN-2001; 2001US-0262321P.

XX (REGC) UNIV CALIFORNIA.

XX Raz E, Lois AF, Takabayashi K;

XX WPI; 2002-062244/08.

XX Modulating cell death or reducing DNA damage in eukaryotic cells, useful
PT for reducing cell death in individual or organ, comprises contacting cell
PT with agent modulating biological activity of DNA-dependent protein
PT kinase.

XX Example 3; Page 33; 57pp; English.

XX The invention relates to a method for modulating cell death or reducing
CC DNA damage in an eukaryotic cell by contacting the cell with an agent
CC that modulates the biological activity of DNA-dependent protein kinase
CC (DNA-PK). The invention also relates nucleic acids which modulate the
CC immune response binding to Ku antigen, resulting in activation of DNA-PK.
CC The method is useful for modulating cell death or reducing DNA damage in
CC an eukaryotic cell, for treating any disorder resulting from a genotoxic
CC insert to a cell e.g., necrosis, apoptosis. The method is also useful for
CC treating cell death-related indications such as Alzheimer's disease,
CC Parkinson's disease, rheumatoid arthritis, septic shock, sepsis, stroke,
CC central nervous system inflammation, osteoporosis, degenerative liver
CC disease, cerebellar degeneration, reperfusion injury, multiple sclerosis,
CC amyotrophic lateral sclerosis, myocardial infarction, head injury damage,
CC acquired immunodeficiency syndrome (AIDS), aplastic anaemia, cerebral
CC infarction, bypass heart surgery, organ transplantation. The method is
CC also useful for treating follicular lymphomas, carcinomas, autoimmune
CC disorders (systemic lupus erythematosus), hormone dependent tumours, The
CC immune mediated glomerulonephritis; apoptosis and viral infections. The
CC present sequence is immunostimulatory oligodeoxynucleotide (ISS-ODN) used
CC for identifying ISS-binding protein, which is used in the exemplification
CC of the invention

XX Sequence 22 BP; 7 A; 2 C; 7 G; 6 T; 0 U; 0 Other;

Query Match 92.7%; Score 20.4; DB 6; Length 22;

Best Local Similarity 95.5%; Pred. No. 3.9;

Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTATAGATGA 22

DB 1 TGACTGTGAACGTTATAGATGA 22

```

RESULT 6
ADO55351
ID ADO55351 standard; DNA; 22 BP.
XX
AC ADO55351;
XX
DT 26-AUG-2004 (first entry)
XX
DE Immune modulatory nucleic acid (IMS) #126.
XX
KW Immune modulatory nucleic acid; IMS; immune modulatory sequence; non CpG;
KW self-molecule related disease; autoimmune disease; multiple sclerosis;
KW rheumatoid arthritis; insulin-dependent diabetes mellitus;
KW autoimmune uveitis; primary biliary cirrhosis; myasthenia gravis;
KW Sjogren's syndrome; pemphigus vulgaris; scleroderma; pernicious anaemia;
KW systemic lupus erythematosus; ankylosing spondylitis;
KW autoimmune skin disease; Grave's disease; inflammatory disease;
KW osteoarthritis; gout; pseudogout; hydroxyapatite deposition disease;
KW asthma; bursitis; tendonitis; conjunctivitis; urethritis; cystitis;
KW balanitis; dermatitis; spinal cord injury; peptic ulcer; hyperlipidaemia;
KW coronary artery disease; migraine; neuroprotective; antirheumatic;
KW antiarthritic; antidiabetic; osteopathic; antigout; antiasthmatic;
KW antiinflammatory; ophthalmological; dermatological; vasotropic;
KW antimigraine; vaccine; gene therapy; ss.
XX
OS Synthetic.
XX
FH Key Location/Qualifiers
FT misc_feature 9..14
FT /tag= a
FT /note= "Core Pu-Pu-X-Y-Py-Py hexamer region"
FT misc_feature 11..12
FT /tag= b
FT /note= "GpG or non-GpG, non-CpG dinucleotide"
XX
FT
XX
PN WO2004047734-A2.
XX
PD 10-JUN-2004.
XX
PF 21-NOV-2003; 2003WO-US037157.
XX
PR 21-NOV-2002; 2002US-0428643P.
XX
XX (BAYH-) BAYHILL THERAPEUTICS INC.
XX (STRD ) UNIV LELAND STANFORD JUNIOR.
XX
PI Garren H, Ho PP, Steinman L;
XX WPI, 2004-441065/41.
XX
XX
XX Pharmaceutical compositions comprising an immune modulatory nucleic acid
XX comprising a hexamer region, useful for treating an autoimmune disease,
XX e.g. multiple sclerosis, rheumatoid arthritis or insulin dependent
XX diabetes mellitus.
XX
XX Example 10; Page 68; 98pp; English.
XX
XX The invention relates to a pharmaceutical composition for treating a
XX disease associated with one or more self-molecules present non-
XX physiologically in an individual (e.g., autoimmune diseases), comprising
XX an immune modulatory nucleic acid (IMS, immune modulatory sequence)
XX comprising a hexamer region of the formula 5'-purine-pyrimidine-[X]-(Y)-
XX pyrimidine-pyrimidine-3', where X and Y are any naturally-occurring or
XX synthetic nucleotides except cytosine-guanine, and a pharmaceutical
XX carrier. The immune modulatory nucleic acid may also contain a polyG
XX region linked 5' and/or 3' to the hexamer region. The invention also
XX relates to a nucleic acid composition comprising a nucleic acid vector
XX having at least one cytosine to non-cytosine substitution (preferably C
XX to G) within a CpG motif, wherein the CpG motif is of the formula: (a) 5'-
XX purine-pyrimidine-C-G-pyrimidine-pyrimidine-3'; or (b) 5'-purine-purine-C
XX -G-pyrimidine-pyrimidine-3'. The immune modulatory nucleic acid sequences
XX are useful in the treatment of disease associated with one or more self-
XX molecules present non-physiologically in an individual, such as

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CC autoimmune diseases (e.g., multiple sclerosis, rheumatoid arthritis,
CC insulin-dependent diabetes mellitus, autoimmune uveitis, primary biliary
CC cirrhosis, myasthenia gravis, Sjogren's syndrome, pemphigus vulgaris,
CC scleroderma, pernicious anaemia, systemic lupus erythematosus, ankylosing
CC spondylitis, autoimmune skin diseases and Grave's disease); inflammatory
CC diseases (e.g., osteoarthritis, gout, pseudogout, hydroxyapatite
CC deposition disease, asthma, bursitis, tendonitis, conjunctivitis,
CC urethritis, cystitis, balanitis and dermatitis); or other conditions such
CC as spinal cord injury, peptic ulcer, hyperlipidaemia, coronary artery
CC disease and migraine. The present sequence represents a specific example
CC of an immune modulatory nucleic acid predicted to be useful for
CC modulating autoimmune disease which is referred to in an example of the
CC invention.
XX
SQ Sequence 22 BP; 7 A; 2 C; 7 G; 6 T; 0 U; 0 Other;
    Query Match          92.7%; Score 20.4; DB 12; Length 22;
    Best Local Similarity 95.5%; Pred. No. 3.9;
    Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Oy 1 TCACCTGTGACGCTTATAGATGA 22
    |||||
Db 1 TGACTGTGACGCTTATAGATGA 22
    |||||
RESULT 7
ADO55287
ID ADO55287 standard; DNA; 22 BP.
XX
AC ADO55287;
XX
DT 26-AUG-2004 (first entry)
XX
DE Immune modulatory nucleic acid (IMS) #62.
XX
KW Immune modulatory nucleic acid; IMS; immune modulatory sequence; non CpG;
KW self-molecule related disease; autoimmune disease; multiple sclerosis;
KW rheumatoid arthritis; insulin-dependent diabetes mellitus;
KW autoimmune uveitis; primary biliary cirrhosis; myasthenia gravis;
KW Sjogren's syndrome; pemphigus vulgaris; scleroderma; pernicious anaemia;
KW systemic lupus erythematosus; ankylosing spondylitis;
KW autoimmune skin disease; Grave's disease; inflammatory disease;
KW osteoarthritis; gout; pseudogout; hydroxyapatite deposition disease;
KW asthma; bursitis; tendonitis; conjunctivitis; urethritis; cystitis;
KW balanitis; dermatitis; spinal cord injury; peptic ulcer; hyperlipidaemia;
KW coronary artery disease; migraine; neuroprotective; antirheumatic;
KW antiarthritic; antidiabetic; osteopathic; antigout; antiasthmatic;
KW antiinflammatory; ophthalmological; dermatological; vasotropic;
KW antimigraine; vaccine; gene therapy; ss.
XX
OS Synthetic.
XX
FH Key Location/Qualifiers
FT misc_feature 9..14
FT /tag= a
FT /note= "Core Pu-Pu-X-Y-Py-Py hexamer region"
FT misc_feature 11..12
FT /tag= b
FT /note= "GpG or non-GpG, non-CpG dinucleotide"
XX
FT modified_base 11
FT /tag= c
FT /mod_base= i
XX
PN WO2004047734-A2.
XX
PD 10-JUN-2004.
XX
PF 21-NOV-2003; 2003WO-US037157.
XX
PR 21-NOV-2002; 2002US-0428643P.
XX (BAYH-) BAYHILL THERAPEUTICS INC.
XX (STRD ) UNIV LELAND STANFORD JUNIOR.

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XX Garren H, Ho PP, Steinman L;
XX WPI; 2004-441065/41.
XX
XX Pharmaceutical compositions comprising an immune modulatory nucleic acid
XX comprising a hexamer region, useful for treating an autoimmune disease,
XX e.g. multiple sclerosis, rheumatoid arthritis or insulin dependent
XX diabetes mellitus.
XX
XX Example 10; Page 66; 98pp; English.
XX
XX The invention relates to a pharmaceutical composition for treating a
XX disease associated with one or more self-molecules present non-
XX physiologically in an individual (e.g., autoimmune diseases), comprising
XX an immune modulatory nucleic acid (IMS, immune modulatory sequence)
XX comprising a hexamer region of the formula 5'-purine-pyrimidine-[X]-[Y]-
XX pyrimidine-pyrimidine-3', where X and Y are any naturally-occurring or
XX synthetic nucleotides except cytosine-guanine, and a pharmaceutical
XX carrier. The immune modulatory nucleic acid may also contain a polyG
XX region linked 5' and/or 3' to the hexamer region. The invention also
XX relates to a nucleic acid composition comprising a nucleic acid vector
XX having at least one cytosine to non-cytosine substitution (preferably C
XX to G) within a CpG motif, wherein the CpG motif is of the formula: (a) 5'-
XX purine-pyrimidine-C-G-pyrimidine-pyrimidine-3'; or (b) 5'-purine-purine-C
XX -G-pyrimidine-pyrimidine-3'. The immune modulatory nucleic acid sequences
XX are useful in the treatment of disease associated with one or more self-
XX molecules present non-physiologically in an individual, such as
XX autoimmune diseases (e.g., multiple sclerosis, rheumatoid arthritis,
XX insulin-dependent diabetes mellitus, autoimmune uveitis, primary biliary
XX cirrhosis, myasthenia gravis, Sjogren's syndrome, pemphigus vulgaris,
XX scleroderma, pernicious anaemia, systemic lupus erythematosus, ankylosing
XX spondylitis, autoimmune skin diseases and Grave's disease); inflammatory
XX diseases (e.g., osteoarthritis, gout, pseudogout, hydroxyapatite
XX deposition disease, asthma, bursitis, tendonitis, conjunctivitis,
XX urethritis, cystitis, balanitis and dermatitis); or other conditions such
XX as spinal cord injury, peptic ulcer, hyperlipidaemia, coronary artery
XX disease and migraine. The present sequence represents a specific example
XX of an immune modulatory nucleic acid predicted to be useful for
XX modulating autoimmune disease which is referred to in an example of the
XX invention.
XX
XX Sequence 22 BP; 7 A; 1 C; 7 G; 6 T; 0 U; 1 Other;
XX
XX Query Match 88.2%; Score 19.4; DB 12; Length 22;
XX Best Local Similarity 90.9%; Pred. No. 12;
XX Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
XX
XX QY 1 TGACTGTGAACGTTATAGATGA 22
XX Db 1 TGACTGTGAACGTTATAGATGA 22
XX
XX RESULT 8
XX AAV32079
XX ID AAV32079 standard; DNA; 22 BP.
XX
XX AC AAV32079;
XX
XX XX 09-SEP-1998 (first entry)
XX
XX DE Nucleotide sequence of DY1018.
XX
XX KW DY1018; beta-gal; ISS-PN/IMM; antigen; immune response; antibody;
XX immunisation; anaphylaxis; IgE; retinopathies; ss.
XX
XX OS Synthetic.
XX
XX Key Location/Qualifiers
XX modified_base 1..22
XX /tag= a
XX /note= "phosphothioate backbone"
XX
XX
XX WO9816247-A1.
XX 23-APR-1998.
XX
XX 09-OCT-1997; 97WO-US019004.
XX
XX 11-OCT-1996; 96US-0028118P.
XX (REGC ) UNIV CALIFORNIA.
XX
XX Carreon DA, Raz B, Roman M;
XX
XX WPI; 1998-261028/23.
XX
XX New immunomodulatory compositions - comprising an antigen conjugated to a
XX polynucleotide that contains an immunostimulatory sequence.
XX
XX Example 1; Page 36; 69pp; English.
XX
XX This is the nucleotide sequence of DY1018, which is conjugated to beta-
XX gal to form ISS-PN/IMM, comprising an immunomodulatory molecule (IMM),
XX which comprises an antigen conjugated to a polynucleotide (PN) that
XX contains at least one immunostimulatory nucleotide sequence (ISS). The
XX conjugate synergistically boost the magnitude of the host immune response
XX against an antigen to a level greater than the host immune response to
XX either the IMM, antigen or ISS-PN alone. These responses to ISS-PN/IMM
XX conjugates are particularly acute during the important early phase of the
XX host immune response to an antigen. The ISS-PN/IMM conjugates boost both
XX humoral (antibody) and cellular (Th1 type) immune responses of the host.
XX Thus, use of the method to boost the immune responsiveness of a host to
XX subsequent challenge by a sensitising antigen without immunisation avoids
XX the risk of Th2-mediated, immunisation-induced anaphylaxis by suppressing
XX IgE production in response to the antigen challenge. The conjugates can
XX also be used to combat pathogenic infection and to stimulate therapeutic
XX angiogenesis to treat conditions in which localised blood flow plays a
XX significant etiological role, e.g. retinopathies
XX
XX Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;
XX
XX Query Match 85.5%; Score 18.8; DB 2; Length 22;
XX Best Local Similarity 90.9%; Pred. No. 24;
XX Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
XX
XX QY 1 TGACTGTGAACGTTATAGATGA 22
XX Db 1 TGACTGTGAACGTTATAGATGA 22
XX
XX RESULT 9
XX AAV80099
XX ID AAV80099 standard; DNA; 22 BP.
XX
XX AC AAV80099;
XX
XX XX 12-MAR-1999 (first entry)
XX
XX DE Immunomodulatory oligo comprising an ISS sequence.
XX
XX KW Immunomodulatory; immunostimulatory; octanucleotide; immune regulation;
XX ISS: cancer; allergy; asthma; hepatitis B infection; papillomavirus;
XX human immunodeficiency virus; influenza; herpes; M. tuberculosis; ss;
XX B. pertussis; malaria; plasmodia; Leishmania; Trypanosoma; Schistosoma.
XX
XX OS Synthetic.
XX
XX WO9855495-A2.
XX
XX 10-DEC-1998.
XX
XX 05-JUN-1998; 98WO-US011578.
XX
XX 06-JUN-1997; 97US-0048793P.
XX

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PA (DYNA-) DYNAVAX TECHNOLOGIES CORP.
XX Schwartz D, Roman M, Dina D;
PI WPI; 1999-059898/05.
XX Immunostimulatory oligonucleotides regulate the immune system - and
PT contain an immune-stimulating octanucleotide sequence; for treating
PT cancer, allergic and infectious diseases.
XX
XX Claim 8; Page 29; 63pp; English.
XX The invention relates to immunomodulatory oligonucleotides that comprise
CC at least 1 immunostimulatory octanucleotide sequence (ISS) where the ISS
CC sequences are selected from the group consisting of AACGTTC, AACGTTCG,
CC GACGTTC, and GACGTTCG. The immunomodulatory sequences are used to treat
CC patients needing immune regulation, such as those suffering from cancer,
CC an allergic disease and asthma. They are also used to prevent infectious
CC diseases such as influenza, herpes, hepatitis B, human immunodeficiency
CC and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and
CC Bordetella pertussis, malarial plasmodia, Leishmania, Trypanosoma and
CC Schistosoma. The immunomodulatory sequences are used to screen for human
CC immunostimulatory activity by incubating macrophage cells and the
CC oligonucleotide; and determining the relative amount of Th1-biased
CC cytokines in the supernatant. Sequences AAV80096 to AAV80103 represent
CC specific claimed examples of such immunomodulatory oligonucleotides
XX
SQ Sequence 22 BP; 6 A; 4 C; 6 G; 6 T; 0 U; 0 Other;
Query Match 85.5%; Score 18.8; DB 2; Length 22;
Best Local Similarity 90.9%; Pred. No. 24;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 TCAGTGTGAACGTTCAGATGA 22
DB 1 TCAGTGTGAACGTTCAGATGA 22

RESULT 10
AAV80097
ID AAV80097 standard; DNA; 22 BP.
XX
XX AAV80097;
XX
XX 12-MAR-1999 (first entry)
XX Immunomodulatory oligo comprising an ISS sequence.
XX Immunomodulatory; immunostimulatory; octanucleotide; immune regulation;
KW ISS: cancer; allergy; asthma; hepatitis B infection; papillomavirus;
KW human immunodeficiency virus; influenza; herpes; M. tuberculosis; ss;
KW B. pertussis; malaria; plasmodia; Leishmania; Trypanosoma; Schistosoma.
XX
XX Synthetic.
XX
XX WO9855495-A2.
XX 10-DEC-1998.
XX
XX 05-JUN-1998; 98WO-US011578.
XX
XX 06-JUN-1997; 97US-0048793P.
XX
XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.
XX Schwartz D, Roman M, Dina D;
XX WPI; 1999-059898/05.
XX Immunostimulatory oligonucleotides regulate the immune system - and
PT contain an immune-stimulating octanucleotide sequence; for treating
PT cancer, allergic and infectious diseases.
XX
XX Claim 24; Page 30; 63pp; English.
XX The invention relates to immunomodulatory oligonucleotides that comprise
CC at least 1 immunostimulatory octanucleotide sequence (ISS) where the ISS
CC sequences are selected from the group consisting of AACGTTC, AACGTTCG,
CC GACGTTC, and GACGTTCG. The immunomodulatory sequences are used to treat
CC patients needing immune regulation, such as those suffering from cancer,
CC an allergic disease and asthma. They are also used to prevent infectious
CC diseases such as influenza, herpes, hepatitis B, human immunodeficiency
CC and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and
CC Bordetella pertussis, malarial plasmodia, Leishmania, Trypanosoma and
CC Schistosoma. The immunomodulatory sequences are used to screen for human
CC immunostimulatory activity by incubating macrophage cells and the
CC oligonucleotide; and determining the relative amount of Th1-biased
CC cytokines in the supernatant. Sequences AAV80096 to AAV80103 represent
CC specific claimed examples of such immunomodulatory oligonucleotides
XX
SQ Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;
Query Match 85.5%; Score 18.8; DB 2; Length 22;
Best Local Similarity 90.9%; Pred. No. 24;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 TCAGTGTGAACGTTCAGATGA 22
DB 1 TCAGTGTGAACGTTCAGATGA 22

RESULT 11
AAV80103
ID AAV80103 standard; DNA; 22 BP.
XX
XX AAV80103;
XX
XX 12-MAR-1999 (first entry)
XX Immunomodulatory oligo comprising an ISS sequence.
XX Immunomodulatory; immunostimulatory; octanucleotide; immune regulation;
KW ISS: cancer; allergy; asthma; hepatitis B infection; papillomavirus;
KW human immunodeficiency virus; influenza; herpes; M. tuberculosis; ss;
KW B. pertussis; malaria; plasmodia; Leishmania; Trypanosoma; Schistosoma.
XX
XX Synthetic.
XX
XX Key Location/Qualifiers
XX modified_base 11
XX /tag= a
XX /note= "5-bromocytosine"
XX
XX WO9855495-A2.
XX 10-DEC-1998.
XX
XX 05-JUN-1998; 98WO-US011578.
XX
XX 06-JUN-1997; 97US-0048793P.
XX
XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.
XX Schwartz D, Roman M, Dina D;
XX WPI; 1999-059898/05.
XX Immunostimulatory oligonucleotides regulate the immune system - and
PT contain an immune-stimulating octanucleotide sequence; for treating
PT cancer, allergic and infectious diseases.
XX
XX Claim 24; Page 30; 63pp; English.
XX The invention relates to immunomodulatory oligonucleotides that comprise
CC at least 1 immunostimulatory octanucleotide sequence (ISS) where the ISS
CC sequences are selected from the group consisting of AACGTTC, AACGTTCG,
CC GACGTTC, and GACGTTCG. The immunomodulatory sequences are used to treat
CC patients needing immune regulation, such as those suffering from cancer,
CC an allergic disease and asthma. They are also used to prevent infectious
CC diseases such as influenza, herpes, hepatitis B, human immunodeficiency
CC and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and
CC Bordetella pertussis, malarial plasmodia, Leishmania, Trypanosoma and
CC Schistosoma. The immunomodulatory sequences are used to screen for human
CC immunostimulatory activity by incubating macrophage cells and the
CC oligonucleotide; and determining the relative amount of Th1-biased
CC cytokines in the supernatant. Sequences AAV80096 to AAV80103 represent
CC specific claimed examples of such immunomodulatory oligonucleotides
XX
SQ Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;
Query Match 85.5%; Score 18.8; DB 2; Length 22;
Best Local Similarity 90.9%; Pred. No. 24;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 TCAGTGTGAACGTTCAGATGA 22
DB 1 TCAGTGTGAACGTTCAGATGA 22

CC GAGGTTCC, and GAGGTTCCG. The immunomodulatory sequences are used to treat
CC patients needing immune regulation, such as those suffering from cancer,
CC an allergic disease and asthma. They are also used to prevent infectious
CC diseases such as influenza, herpes, hepatitis B, human immunodeficiency
CC and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and
CC Bordetella pertussis, malarial plasmodia, Leishmania, Trypanosoma and
CC Schistosoma. The immunomodulatory sequences are used to screen for human
CC immunostimulatory activity by incubating macrophage cells and the
CC oligonucleotide; and determining the relative amount of Th1-biased
CC cytokines in the supernatant. Sequences AAV80096 to AAV80103 represent
CC specific claimed examples of such immunomodulatory oligonucleotides

XX
SQ Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;

Query Match 85.5%; Score 18.8; DB 2; Length 22;
Best Local Similarity 90.9%; Pred. No. 24;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 TGACTGTGACGTTATAGATCA 22
Db 1 TGACTGTGACGTTGCGATGA 22

RESULT 12
AAV80106
ID AAV80106 standard; DNA; 22 BP.

XX AAV80106;

XX 12-MAR-1999 (first entry)

DE Oligo used in experiments for stimulation of cytokine production.

XX Immunomodulatory; immunostimulatory; octanucleotide; immune regulation;
KW ISS: cancer; allergy; asthma; hepatitis B infection; papillomavirus;
KW human immunodeficiency virus; influenza; herpes; M. tuberculosis; ss;
KW B. pertussis; malaria; plasmodia; leishmania; Trypanosoma; Schistosoma.

XX Synthetic.

XX WO9855495-A2.

XX 10-DEC-1998.

XX 05-JUN-1998; 98WO-US011578.

XX 06-JUN-1997; 97US-0048793P.

XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.

XX Schwartz D, Roman M, Dina D;

XX WPI; 1999-059898/05.

XX Immunostimulatory oligonucleotides regulate the immune system - and
XX contain an immune-stimulating octanucleotide sequence; for treating
XX cancer, allergic and infectious diseases.

XX Example 1; Page 29; 63pp; English.

XX The invention relates to immunomodulatory oligonucleotides that comprise
XX at least 1 immunostimulatory octanucleotide sequence (ISS) where the ISS
XX sequences are selected from the group consisting of AACGTTCC, AACGTTCCG,
XX GAGGTTCC, and GAGGTTCCG. The immunomodulatory sequences are used to treat
XX patients needing immune regulation, such as those suffering from cancer,
XX an allergic disease and asthma. They are also used to prevent infectious
XX diseases such as influenza, herpes, hepatitis B, human immunodeficiency
XX and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and
XX Bordetella pertussis, malarial plasmodia, Leishmania, Trypanosoma and
XX Schistosoma. The immunomodulatory sequences are used to screen for human
XX immunostimulatory activity by incubating macrophage cells and the
XX oligonucleotide; and determining the relative amount of Th1-biased
XX cytokines in the supernatant. Sequences AAV80104 to AAV80116 represent

CC oligonucleotides that were tested for immunostimulatory activity. These
CC were used in experiments for the stimulation of cytokine production and
CC were found to lack immunostimulatory activity. The invention provides
CC specific claimed examples (AAV80096-103) of immunomodulatory sequences

XX
SQ Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;

Query Match 85.5%; Score 18.8; DB 2; Length 22;
Best Local Similarity 90.9%; Pred. No. 24;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 TGACTGTGACGTTATAGATCA 22
Db 1 TGACTGTGACGTTGCGATGA 22

RESULT 13
AAV80101
ID AAV80101 standard; DNA; 22 BP.

XX AAV80101;

XX 12-MAR-1999 (first entry)

DE Immunomodulatory oligo comprising an ISS sequence.

XX Immunomodulatory; immunostimulatory; octanucleotide; immune regulation;
KW ISS: cancer; allergy; asthma; hepatitis B infection; papillomavirus;
KW human immunodeficiency virus; influenza; herpes; M. tuberculosis; ss;
KW B. pertussis; malaria; plasmodia; leishmania; Trypanosoma; Schistosoma.

XX Synthetic.

XX Key Location/Qualifiers

XX modified_base 11
XX /*tag= a
XX /note= "5-bromocytosine"

XX WO9855495-A2.

XX 10-DEC-1998.

XX 05-JUN-1998; 98WO-US011578.

XX 06-JUN-1997; 97US-0048793P.

XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.

XX Schwartz D, Roman M, Dina D;

XX WPI; 1999-059898/05.

XX Immunostimulatory oligonucleotides regulate the immune system - and
XX contain an immune-stimulating octanucleotide sequence; for treating
XX cancer, allergic and infectious diseases.

XX Claim 22; Page 30; 63pp; English.

XX The invention relates to immunomodulatory oligonucleotides that comprise
XX at least 1 immunostimulatory octanucleotide sequence (ISS) where the ISS
XX sequences are selected from the group consisting of AACGTTCC, AACGTTCCG,
XX GAGGTTCC, and GAGGTTCCG. The immunomodulatory sequences are used to treat
XX patients needing immune regulation, such as those suffering from cancer,
XX an allergic disease and asthma. They are also used to prevent infectious
XX diseases such as influenza, herpes, hepatitis B, human immunodeficiency
XX and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and
XX Bordetella pertussis, malarial plasmodia, Leishmania, Trypanosoma and
XX Schistosoma. The immunomodulatory sequences are used to screen for human
XX immunostimulatory activity by incubating macrophage cells and the
XX oligonucleotide; and determining the relative amount of Th1-biased
XX cytokines in the supernatant. Sequences AAV80096 to AAV80103 represent
XX specific claimed examples of such immunomodulatory oligonucleotides

```
SQ Sequence 22 BP; 6 A; 4 C; 6 G; 6 T; 0 U; 0 Other;
  Query Match      85.5%; Score 18.8; DB 2; Length 22;
  Best Local Similarity 90.9%; Pred. No. 24;
  Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTATAGATGA 22
Db 1 TGACTGTGAACGTTCCAGATGA 22

RESULT 14
AAV80104
ID AAV80104 standard; DNA; 22 BP.
XX
AC AAV80104;
XX
DT 12-MAR-1999 (first entry)
XX
DE Oligo used in experiments for stimulation of cytokine production.
XX
KW Immunomodulatory; immunostimulatory; octanucleotide; immune regulation;
KW ISS: cancer; allergy; asthma; hepatitis B infection; papillomavirus;
KW human immunodeficiency virus; influenza; herpes; M. tuberculosis; ss;
KW B. pertussis; malaria; plasmodia; leishmania; trypanosoma; schistosoma.
XX
OS Synthetic.
XX
PN WO9855495-A2.
XX
PD 10-DEC-1998.
XX
PF 05-JUN-1998; 98WO-US011578.
XX
PR 06-JUN-1997; 97US-0048793P.
XX
PA (DYNA-) DYNAVAX TECHNOLOGIES CORP.
XX
PI Schwartz D, Roman M, Dina D;
XX
WPI; 1999-059898/05.
XX
DR Immunostimulatory oligonucleotides regulate the immune system - and
XX contain an immune-stimulating octanucleotide sequence; for treating
XX cancer, allergic and infectious diseases.
XX
PS Example 1; Page 29; 63pp; English.
XX
CC The invention relates to immunomodulatory oligonucleotides that comprise
CC at least 1 immunostimulatory octanucleotide sequence (ISS) where the ISS
CC sequences are selected from the group consisting of AACGTTC, AACGTTCG,
CC GACGTTC, and GACGTTCG. The immunomodulatory sequences are used to treat
CC patients needing immune regulation, such as those suffering from cancer,
CC an allergic disease and asthma. They are also used to prevent infectious
CC diseases such as influenza, herpes, hepatitis B, human immunodeficiency
CC disease, and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and
CC Bordetella pertussis, malarial plasmodia, Leishmania, Trypanosoma and
CC Schistosoma. The immunomodulatory sequences are used to screen for human
CC immunostimulatory activity by incubating macrophage cells and the
CC oligonucleotide; and determining the relative amount of Th1-biased
CC cytokines in the supernatant. Sequences AAV80104 to AAV80116 represent
CC oligonucleotides that were tested for immunostimulatory activity. These
CC were used in experiments for the stimulation of cytokine production and
CC were found to lack immunostimulatory activity. The invention provides
CC specific claimed examples (AAV80096-103) of immunomodulatory sequences
XX
SQ Sequence 22 BP; 7 A; 1 C; 8 G; 6 T; 0 U; 0 Other;
  Query Match      85.5%; Score 18.8; DB 2; Length 22;
  Best Local Similarity 90.9%; Pred. No. 24;
  Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTATAGATGA 22
```

```
Db 1 TGACTGTGAAGTTAGATGA 22

RESULT 15
AAV80102
ID AAV80102 standard; DNA; 22 BP.
XX
AC AAV80102;
XX
DT 12-MAR-1999 (first entry)
XX
DE Immunomodulatory oligo comprising an ISS sequence.
XX
KW Immunomodulatory; immunostimulatory; octanucleotide; immune regulation;
KW ISS: cancer; allergy; asthma; hepatitis B infection; papillomavirus;
KW human immunodeficiency virus; influenza; herpes; M. tuberculosis; ss;
KW B. pertussis; malaria; plasmodia; leishmania; trypanosoma; schistosoma.
XX
OS Synthetic.
XX
Key Location/Qualifiers
FH modified_base 11
FT /*tag= a
FT /note= "5-bromocytosine"
XX
PN WO9855495-A2.
XX
PD 10-DEC-1998.
XX
PF 05-JUN-1998; 98WO-US011578.
XX
PR 06-JUN-1997; 97US-0048793P.
XX
PA (DYNA-) DYNAVAX TECHNOLOGIES CORP.
XX
PI Schwartz D, Roman M, Dina D;
XX
WPI; 1999-059898/05.
XX
DR Immunostimulatory oligonucleotides regulate the immune system - and
XX contain an immune-stimulating octanucleotide sequence; for treating
XX cancer, allergic and infectious diseases.
XX
PS Claim 23; Page 30; 63pp; English.
XX
CC The invention relates to immunomodulatory oligonucleotides that comprise
CC at least 1 immunostimulatory octanucleotide sequence (ISS) where the ISS
CC sequences are selected from the group consisting of AACGTTC, AACGTTCG,
CC GACGTTC, and GACGTTCG. The immunomodulatory sequences are used to treat
CC patients needing immune regulation, such as those suffering from cancer,
CC an allergic disease and asthma. They are also used to prevent infectious
CC diseases such as influenza, herpes, hepatitis B, human immunodeficiency
CC disease, and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and
CC Bordetella pertussis, malarial plasmodia, Leishmania, Trypanosoma and
CC Schistosoma. The immunomodulatory sequences are used to screen for human
CC immunostimulatory activity by incubating macrophage cells and the
CC oligonucleotide; and determining the relative amount of Th1-biased
CC cytokines in the supernatant. Sequences AAV80096 to AAV80103 represent
CC specific claimed examples of such immunomodulatory oligonucleotides
XX
SQ Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;
  Query Match      85.5%; Score 18.8; DB 2; Length 22;
  Best Local Similarity 90.9%; Pred. No. 24;
  Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTATAGATGA 22
Db 1 TGACTGTGAACGTTCCAGATGA 22

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